**DATE:** Sept. 03, 2009

**SUBJECT:** Chemicals Evaluated for Carcinogenic Potential by the Office of Pesticide Programs

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TO: Division Directors AD, BPPD, EFED, FEAD, HED, RD and SRRD

The attached list provides an overview of chemicals evaluated for carcinogenic potential by the Health Effects Division (HED) of the Office of Pesticide Programs (OPP) through August 2009. Applying the Agency's Guidelines for Carcinogen Risk Assessment, the classification of the chemical is made by HED's Cancer Assessment Review Committee (CARC) or, in the case of where there is no evidence of carcinogenicity, by the HED Risk Assessment Team.

This list includes the chemical name, CAS Number, PC code, the cancer classification, report date, species, tumor types, and, if required, the human equivalency potency factor (Q1\*). The potency factor (Q1\*), unless otherwise indicated, is based on the oral route. The Q1\* is expressed as (mg/kg/day)-1 for the oral route and as (mg/m3)-1 for the inhalation route.

It should be noted that the evaluation of many of these chemicals is an ongoing process, therefore, the information in this list (i.e., classification and/or the quantification) may be subject to change as new and/or additional data are submitted to OPP. This list should not be used as the single source for either the classification or quantification of the carcinogenic potential. This list will be updated annually.

If further information is required please contact me (Phone: 703-308-6175; E-mail: may.brenda@epa.gov).

Science Information Management Branch
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#### **BACKGROUND**

#### What is this list?

The Chemicals Evaluated for Carcinogenic Potential provides an overview of the compounds evaluated for carcinogenicity by the Health Effects Division of the Office of Pesticide Programs.

**NOTE:** As new information becomes available, the list may become out-of-date. Therefore, it should not be used as the sole reference regarding the carcinogenic potential for a pesticide. EPA intends to update the list each year to include new evaluations or re-evaluations.

#### How does EPA review pesticides for potential carcinogenicity?

The Health Effects Division of the Office of Pesticide Programs performs an independent review of studies conducted in mice and rats to evaluate the carcinogenic potential of pesticides. The results of the independent review are peer-reviewed by the Cancer Assessment Review Committee. This committee recommends a cancer classification. The classification will determine how the Agency regulates the pesticide and will include methods for quantification of human risk. In some cases, EPA also requests review by the FIFRA Scientific Advisory Panel.

#### What factors does EPA consider in its review of cancer risk?

When assessing possible cancer risk posed by a pesticide, EPA considers how strongly carcinogenic the chemical is (its potency) and the potential for human exposure. The pesticides are evaluated not only to determine if they cause cancer in laboratory animals, but also as to their potential to cause human cancer. For any pesticide classified as a potential carcinogen, the risk would depend on the extent to which a person might be exposed (how much time and to what quantity of the pesticide). The factors considered include short-term studies, long-term cancer studies, mutagenicity studies, and structure activity concerns. (The term "weight-of-the-evidence" is used in referring to such a review. This means that the recommendation is not based on the results of one study, but on the results of all studies that are available.)

#### When does EPA review pesticides for potential carcinogenicity?

EPA reviews studies submitted when a pesticide is proposed for registration. Studies are required in two species (mice and rats) and two sexes (males and females). These studies are required for all pesticides used on food and some non-food pesticides that could lead to long-term exposures in humans. These studies may be reviewed again when a pesticide undergoes reregistration and the cancer classification may be reevaluated, particularly if new studies have been submitted.

#### Why are there several different cancer classifications in the list?

EPA's guidelines for evaluating the potential carcinogenicity of chemicals have been updated over the years to reflect increased understanding of ways chemicals may cause cancer. The current guidelines call for greater emphasis on characterization discussions for hazard, doseresponse assessment, exposure assessment, and risk characterization, as well as the use of mode of action in the assessment of potential carcinogenesis.

EPA does not have the resources to re-evaluate every chemical to determine how it would be described under new guidelines, and there is no reason to re-evaluate chemicals unless there is some new information that could change the basic understanding of that chemical.

#### How have the guidelines changed?

EPA issued its first set of principles to guide evaluation of human cancer potential in1976. In 1986, EPA issued updated guidance, which included a letter system (A-E) for designating degree of carcinogenic potential. In the 1986 guidelines, hazard identification and the weight-of evidence process focused on tumor findings. The human carcinogenic potential of agents was characterized by a six-category alphanumeric classification system (A, B1, B2, C, and D). In 1996, EPA released "Proposed Guidelines for Carcinogen Risk Assessment," which used descriptive phrases rather than the alphanumeric classification to classify carcinogenic potential. In the 1996 classification structure, increased emphasis was placed on discussing characterization of hazard, dose-response, and exposure assessments. The hazard and weight of evidence process embraced an analysis of all relevant biological information and emphasized understanding the agent's mode of action in producing tumors to reduce the uncertainty in describing the likelihood of harm. By 1999, the science related to carcinogens had advanced significantly. EPA issued draft guidelines that continued the greater emphasis on characterization discussions for hazard, dose-response assessment, exposure assessment, risk characterization and the use of mode of action in the assessment of potential carcinogenesis. In addition, the guidelines included consideration of risk to children, as well as addressing other issues such as nuances related to the amount and adequacy of data on a chemical.

In March, 2005, EPA released its final *Guidelines for Carcinogen Risk Assessment* (EPA/630/P-03/001B). These guidelines represent the culmination of a long development process, replacing EPA's original cancer risk assessment guidelines (1986) and its interim final guidelines (1999). http://www.epa.gov/cancerguidelines/

#### How do the different designations compare?

The short answer is that they cannot be directly compared. Each system designation refers to the reviews and criteria it contains. A substance that is, for example, a "C" in the 1986 system may not be directly translatable to any particular category in the later systems. The designation for any substance must be considered in the context of the system under which it was reviewed.

A list of the descriptors from the various classification systems and their definitions are given on the following pages.

# Carcinogenicity Classification of Pesticides: Derivation and Definition of Terms

#### **CLASSIFICATION-2005**

The following descriptors from the 2005 Guidelines for Carcinogen Risk Assessment can be used as an introduction to the weight of evidence narrative in the cancer risk assessment. The examples presented in the discussion of the descriptors are illustrative. The examples are neither a checklist nor a limitation for the descriptor. The complete weight of evidence narrative, rather than the descriptor alone, provides the conclusions and the basis for them.

**CARCINOGENIC TO HUMANS.** This descriptor indicates strong evidence of human carcinogenicity. It covers different combinations of evidence.

- This descriptor is appropriate when there is convincing epidemiologic evidence of a causal association between human exposure and cancer.
- Exceptionally, this descriptor may be equally appropriate with a lesser weight of epidemiologic evidence that is strengthened by other lines of evidence. It can be used when all of the following conditions are met: (a) there is strong evidence of an association between human exposure and either cancer or the key precursor events of the agent's mode of action but not enough for a causal association, and (b) there is extensive evidence of carcinogenicity in animals, and (c) the mode(s) of carcinogenic action and associated key precursor events have been identified in animals, and (d) there is strong evidence that the key precursor events that precede the cancer response in animals are anticipated to occur in humans and progress to tumors, based on available biological information. In this case, the narrative includes a summary of both the experimental and epidemiologic information on mode of action and also an indication of the relative weight that each source of information carries, e.g., based on human information, and based on limited human and extensive animal experiments.

**LIKELY TO BE CARCINOGENIC TO HUMANS.** This descriptor is appropriate when the weight of the evidence is adequate to demonstrate carcinogenic potential to humans but does not reach the weight of evidence for the descriptor "Carcinogenic to Humans." Adequate evidence consistent with this descriptor covers a broad spectrum. As stated previously, the use of the term "likely" as a weight of evidence descriptor does not correspond to a quantifiable probability. The examples below are meant to represent the broad range of data combinations that are covered by this descriptor; they are illustrative and provide neither a checklist nor a limitation for the data that might support use of this descriptor.

Moreover, additional information, e.g., on mode of action, might change the choice of descriptor for the illustrated examples. Supporting data for this descriptor may include:

 an agent demonstrating a plausible (but not definitively causal) association between human exposure and cancer, in most cases with some supporting biological, experimental evidence, though not necessarily carcinogenicity data from animal experiments;

- an agent that has tested positive in animal experiments in more than one species, sex, strain, site, or exposure route, with or without evidence of carcinogenicity in humans;
- a positive tumor study that raises additional biological concerns beyond that of a statistically significant result, for example, a high degree of malignancy, or an early age at onset;
- a rare animal tumor response in a single experiment that is assumed to be relevant to humans; or
- a positive tumor study that is strengthened by other lines of evidence, for example, either plausible (but not definitively causal) association between human exposure and cancer or evidence that the agent or an important metabolite causes events generally known to be associated with tumor formation (such as DNA reactivity or effects on cell growth control) likely to be related to the tumor response in this case.

**SUGGESTIVE EVIDENCE OF CARCINOGENIC POTENTIAL.** This descriptor of the database is appropriate when the weight of evidence is suggestive of carcinogenicity; a concern for potential carcinogenic effects in humans is raised, but the data are judged not sufficient for a stronger conclusion. This descriptor covers a spectrum of evidence associated with varying levels of concern for carcinogenicity, ranging from a positive cancer result in the only study on an agent to a single positive cancer result in an extensive database that includes negative studies in other species. Depending on the extent of the database, additional studies may or may not provide further insights. Some examples include:

- a small, and possibly not statistically significant, increase in tumor incidence observed in a single animal or human study that does not
  reach the weight of evidence for the descriptor "Likely to Be Carcinogenic to Humans." The study generally would not be contradicted by
  other studies of equal quality in the same population group or experimental system (see discussions of conflicting evidence and differing
  results, below);
- a small increase in a tumor with a high background rate in that sex and strain, when there is some but insufficient evidence that the observed tumors may be due to intrinsic factors that cause background tumors and not due to the agent being assessed. (When there is a high background rate of a specific tumor in animals of a particular sex and strain, then there may be biological factors operating independently of the agent being assessed that could be responsible for the development of the observed tumors.) In this case, the reasons for determining that the tumors are not due to the agent are explained;
- evidence of a positive response in a study whose power, design, or conduct limits the ability to draw a confident conclusion (but does not
  make the study fatally flawed), but where the carcinogenic potential is strengthened by other lines of evidence (such as structure-activity
  relationships); or
- a statistically significant increase at one dose only, but no significant response at the other doses and no overall trend.

**INADEQUATE INFORMATION TO ASSESS CARCINOGENIC POTENTIAL.** This descriptor of the database is appropriate when available data are judged inadequate for applying one of the other descriptors. Additional studies generally would be expected to provide further insights. Some examples include:

- little or no pertinent information;
- conflicting evidence, that is, some studies provide evidence of carcinogenicity but other studies of equal quality in the same sex and strain are negative. Differing results, that is, positive results in some studies and negative results in one or more different experimental

- systems, do not constitute *conflicting evidence*, as the term is used here. Depending on the overall weight of evidence, differing results can be considered either suggestive evidence or likely evidence; or
- negative results that are not sufficiently robust for the descriptor, "Not Likely to Be Carcinogenic to Humans."

**NOT LIKELY TO BE CARCINOGENIC TO HUMANS.** This descriptor is appropriate when the available data are considered robust for deciding that there is no basis for human hazard concern. In some instances, there can be positive results in experimental animals when there is strong, consistent evidence that each mode of action in experimental animals does not operate in humans. In other cases, there can be convincing evidence in both humans and animals that the agent is not carcinogenic. The judgment may be based on data such as:

- animal evidence that demonstrates lack of carcinogenic effect in both sexes in well-designed and well-conducted studies in at least two
  appropriate animal species (in the absence of other animal or human data suggesting a potential for cancer effects),
- convincing and extensive experimental evidence showing that the only carcinogenic effects observed in animals are not relevant to humans,
- convincing evidence that carcinogenic effects are not likely by a particular exposure route (see Section 2.3), or
- convincing evidence that carcinogenic effects are not likely below a defined dose range.

A descriptor of "not likely" applies only to the circumstances supported by the data. For example, an agent may be "Not Likely to Be Carcinogenic" by one route but not necessarily by another. In those cases that have positive animal experiment(s) but the results are judged to be not relevant to humans, the narrative discusses why the results are not relevant.

**MULTIPLE DESCRIPTORS.** More than one descriptor can be used when an agent's effects differ by dose or exposure route. For example, an agent may be "Carcinogenic to Humans" by one exposure route but "Not Likely to Be Carcinogenic" by a route by which it is not absorbed. Also, an agent could be "Likely to Be Carcinogenic" above a specified dose but "Not Likely to Be Carcinogenic" below that dose because a key event in tumor formation does not occur below that dose.

#### **CLASSIFICATION -1999 Draft**

The terms used to describe carcinogenic potential in the July 1999 "Review Draft of the Guidelines for Carcinogen Risk Assessment" are listed and defined as follows:

**CARCINOGENIC TO HUMANS**. This descriptor is appropriate when there is convincing epidemiologic evidence demonstrating causality between human exposure and cancer. This descriptor is also appropriate when there is an absence of conclusive epidemiologic evidence to clearly establish a cause and effect relationship between human exposure and cancer, but there is compelling evidence of carcinogenicity in animals and mechanistic information in animals and humans demonstrating similar mode(s) of carcinogenic action. It is used when all of the following conditions are met:

- There is evidence in a human population(s) of association of exposure to the agent with cancer, but not enough to show a causal association, and
- There is extensive evidence of carcinogenicity, and
- The mode(s) of carcinogenic action and associated key events have been identified in animals, and
- The keys events that precede the cancer response in animals have been observed in the human population(s) that also shows evidence of an association of exposure to the agent with cancer.

**LIKELY TO BE CARCINOGENIC TO HUMANS.** This descriptor is appropriate when the available tumor effects and other key data are adequate to demonstrate carcinogenic potential to humans. Adequate data are within a spectrum. At one end is evidence for an association between human exposure to the agent and cancer and strong experimental evidence of carcinogenicity in animals; at the other, with no human data, the weight of experimental evidence shows animal carcinogenicity by a mode or modes of action that are relevant or assumed to be relevant to humans.

SUGGESTIVE EVIDENCE OF CARCINOGENICITY, BUT NOT SUFFICIENT TO ASSESS HUMAN CARCINOGENIC POTENTIAL. This descriptor is appropriate when the evidence from human or animal data is suggestive of carcinogenicity, which raises a concern for carcinogenic effects but is judged not sufficient for a conclusion as to human carcinogenic potential. Examples of such evidence may include: a marginal increase in tumors that may be exposure-related, or evidence is observed only in a single study, or the only evidence is limited to certain high background tumors in one sex of one species. Dose-response assessment is not indicated for these agents. Further studies would be needed to determine human carcinogenic potential.

**DATA ARE INADEQUATE FOR AN ASSESSMENT OF HUMAN CARCINOGENIC POTENTIAL**. This descriptor is used when available data are judged inadequate to perform an assessment. This includes a case when there is a lack of pertinent or useful data or when existing evidence is conflicting, e.g., some evidence is suggestive of carcinogenic effects, but other equally pertinent evidence does not confirm a concern.

**NOT LIKELY TO BE CARCINOGENIC TO HUMANS**. This descriptor is used when the available data are considered robust for deciding that there is no basis for human hazard concern. The judgment may be based on:

- Extensive human experience that demonstrates lack of carcinogenic effect (e.g., phenobarbital).
- Animal evidence that demonstrates lack of carcinogenic effect in at least two well- designed and well-conducted studies in two
  appropriate animal species (in the absence of human data suggesting a potential for cancer effects).
- Extensive experimental evidence showing that the only carcinogenic effects observed in animals are not considered relevant to humans (e.g., showing only effects in the male rat kidney due to accumulation of alpha<sub>2u</sub>-globulin).
- Evidence that carcinogenic effects are not likely by a particular route of exposure.
- Evidence that carcinogenic effects are not anticipated below a defined dose range.

#### **CLASSIFICATION-1996**

In April 1996, EPA released the "Proposed Guidelines for Carcinogen Risk Assessment." This scheme varied from the earlier 1986 scheme in that it used descriptors rather than letters to classify carcinogenic potential. The descriptors are:

**KNOWN/LIKELY**. This category of descriptors is appropriate when the available tumor effects and other key data are adequate to convincingly demonstrate carcinogenic potential for humans.

**CANNOT BE DETERMINED**. This category of descriptors is appropriate when available tumor effects or other key data are suggestive or conflicting or limited in quantity and, thus, are not adequate to convincingly demonstrate carcinogenic potential for humans. In general, further agent specific and generic research and testing are needed to be able to describe human carcinogenic potential.

**NOT LIKELY**. This is the appropriate descriptor when experimental evidence is satisfactory for deciding that there is no basis for human hazard concern, as follows (in the absence of human data suggesting a potential for cancer effects).

#### **CLASSIFICATION -1986**

The following cancer classification scheme was first introduced in 1986. It was used until 1996.

**GROUP A-HUMAN CARCINOGEN**. This group is used only when there is sufficient evidence from epidemiologic studies to support a causal association between exposure to the agents and cancer.

**GROUP B-PROBABLE HUMAN CARCINOGEN**. This group includes agents for which the weight of evidence of human carcinogenicity based on epidemiologic studies is "limited" and also includes agents for which the weight of evidence of carcinogenicity based on animal studies is "sufficient." The group is divided into two subgroups. **Group B1** is reserved for agents for which there is limited evidence of

carcinogenicity from epidemiologic studies. **Group B2** is used for Agents for which there is "sufficient: evidence from animal studies and for which there is "inadequate evidence" or "no data" from epidemiologic studies.

**GROUP C-POSSIBLE HUMAN CARCINOGEN**. This group is used for agents with limited evidence of carcinogenicity in animals in the absence of human data.

**GROUP D-NOT CLASSIFIABLE AS TO HUMAN CARCINOGENICITY**. This group is generally used for agents with inadequate human and animal evidence of carcinogenicity or for which no data are available.

**GROUP E-EVIDENCE OF NON-CARCINOGENICITY FOR HUMANS**. This group is used for agents that show no evidence for carcinogenicity in at least two adequate animal tests in different species or in both adequate epidemiologic and animal studies.

#### OTHER DEFINITIONS

#### Quantification of Cancer Risk - Carcinogenic Potency Factor (Q<sub>1</sub>\*)

Q1 STAR (Q1\*) - In the classification of human or probable-human carcinogens, mathematical models are used to estimate an upper-bound excess cancer risk associated with lifetime ingestion in the diet. The data used in these estimates usually come from lifetime exposure studies in animals. The USEPA generally uses the linearized multistage model for its cancer risk assessment. This model fits linear dose-response curves to low doses and is consistent with a no-threshold model of carcinogenesis, i.e., exposure to even a very small amount of the substance produces a finite increased risk of cancer.

The linearized multistage model uses dose-response data from the most appropriate carcinogenic study to calculate a carcinogenic potency factor (q<sub>1</sub>\*) for humans. The q<sub>1</sub>\* is then used to determine the concentrations of the chemical in the diet that are associated with theoretical upperbound excess lifetime cancer risks of 1 in 10,000, 1 in 100,000, and 1 in 1,000,000 (10-4, 10-5, 10-6 respectively) individuals over a lifetime of exposure.

**Mode of Action (MOA)** - The key cellular and biochemical events that have to happen for a biological effect to develop. Mode of action is contrasted with mechanism of action which is a more complete understanding of the step by step pathway leading to a biological effect. Some established MOAs include:

**Androgen Dependent -** The chemical disrupts the normal levels of reproductive hormones (e.g., testosterone, luteinizing hormone) which in turn stimulates the target tissue (e.g., leydig cells, testicular tissue) to divide which may lead to hyperplasia and neoplasia. For agents to pose a hazard to humans by this MOA, sufficient exposure levels need to be encountered which produce the same level of biological effect as seen in rodents. This is consistent with the MOA for Leydig cell tumorigenesis.

**Cytotoxicity and Regenerative Proliferation -** Continuous exposure to a chemical or its metabolite causes persistent cell killing which in turn may result in a persistent regenerative proliferative response in the damaged tissue. For irreversible tissue alterations to occur in humans, including cancer by this mode of action, a sufficient exposure must be encountered over a prolonged period.

**Mitogenesis -** Mitogenic chemicals act by promoting the clonal expansion of preneoplastic cells by stimulating cell proliferation. This mode of action is frequently found in the rodent liver where it is generally associated with an increase in metabolizing enzymes. A mitogenic chemical stimulates cell proliferation in the target organ without obvious cytotoxicity or cell death. Another important feature of this MOA is that the mitogenic effect is not persistent over time; instead it is resolved and then is manifested within proliferative foci which are considered preneoplastic lesions. Through continuous exposure, it is these preneoplastic lesions that develop into tumors. At this time, the adverse health effects caused by this MOA are presumed to be relevant to humans.

**Mutagenesis** - The chemical or a metabolite has the ability to react with or bind DNA in a manner that causes mutations. It is usually positive in multiple test systems for different genetic endpoints (particularly gene mutations and structural chromosome aberrations) and in tests performed *in vivo* and *in vitro*. Adverse health effects in rodents from these chemicals are considered relevant for human health risk.

**Neuroendrocrine Disruption -** Chemicals that disrupt hypothalamic control of pituitary function leading to a decrease in hormone release (e.g., luteinizing hormone) and the disruption of the ovarian cycle. This may result in an increase in cell proliferation in the mammary gland due to a hyperstimulation by estrogen. In the case of chloro-s-triazines, this neuroendocrine MOA is not considered relevant to humans because it depends on a rodent specific reproductive process.

**PPAR-alpha Agonism -** Chemicals that bind to and activate the Peroxisome Proliferator-Activated Receptor (PPAR) stimulate biological responses in the liver (e.g., peroxisome proliferation, induction of lipid metabolizing enzymes, oxidative stress, and hepatocyte mitogenesis). Activation of PPAR-alpha results in an increase in cell proliferation and clonal expansion of preneoplastic foci in the liver. While the human relevance of this MOA has not been definitively determined, most of the evidence indicates that this mode of action is not operative in the human liver.

**Thyroid Hormone Disruption -** Disruption of normal levels of thyroid hormones may lead to an increase of thyroid stimulating hormone (TSH) which results in an increase in cell proliferation of the thyroid gland. If exposure is continuous in the animal, thyroid follicular cell tumors can potentially develop. However, the development of thyroid cancer by this mode of action in humans is considered unlikely since prolonged stimulation of the thyroid gland by TSH has not been associated with tumorigenesis in humans. However, this MOA is relevant as an indicator for potential noncancer health effects (e.g., goiter, neurodevelopmental, etc) due thyroid disruption in humans.

CAS NO.	PC CODE	CANCER CLASSIFICATION		QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
		Not Likely To Be Carcinogenic	DATE	METHOD	
89415-87-2	128826		8/28/2000	NR	Not Applicable
00110012			0,20,200		- Total Application
2758-42-1	030819	To Humans	6/13/2003	NR	Not Applicable
		Not Likely To Be Carcinogenic			
94-82-6	030801	To Humans	6/13/2003	NR	Not Applicable
		Group DNot Classifiable as to			
94-75-7	030001	Human Carcinogenicity	1/29/1997	NR	Not Applicable
		Not Likely To Be Carcinogenic			
15165-67-0	031402	To Humans	8/13/2007	NR	Not Applicable
		Group CPossible Human			Kidney tumors in B6C3F1 mice (M)
120-32-1	062201	Carcinogen	9/5/1995	RfD Approach	Kidney tumors in F344/N rats (F)
		Group DNot Classifiable As			
504-24-5	069201	To Human Carcinogenicity	8/6/2007	NR	Not Applicable
		Group CPossible Human			
30560-19-1	103301	Carcinogen	5/8/1985	NR	Liver tumors in CD-1 mice (F)
		Nat Libert Ta Da Canaina mania			
F7000 40 7	000000	,	44/40/0000	ND	Not A - Post I
57960-19-7	006329		11/13/2003	NK	Not Applicable
00.05.5	444404		F/00/4000	ND	Liver tumors in Wistar rats (M)
60-35-5	111101	Carcinogen	5/29/1990	NK	Liver tumors in F344 rats (M & F)
		Not Likely To Do Carsing renie			
125/110 20 7	000050		12/11/2001	ND	Not Applicable
135410-20-7	099050	TO Humans	12/11/2001	INIX	Lung tumors in CD- 1 mice (M & F)
					Ovarian tumors in CD-1 mice (W & F)  Stablished a cytotoxic
					(secondary to oxidative damage by a reactive quinone imine
		Suggestive Evidence of			intermediate) mode of action for the nasal olfactory epithelial
3/256-82-1	121601		1/3/2007	PfD Approach	tumors and a hormonal mode of action for
34230-62-1	121001		1/3/2007	Кір Арріоасіі	tumors and a normonal mode of action for
135158-54-2	061402		12/9/1999	NR	Not Applicable
100100-04-2	001702		12/3/1333	INIX	140t / Applicable
					Liver tumors in B6C3F1 (M & F)
					Liver tumors in CD-1 mice (M & F); Established a PPARa
62476-59-9	114402		7/9/2003	MOF Approach	mode of action for liver tumors in mice
	89415-87-2 2758-42-1 94-82-6 94-75-7 15165-67-0 120-32-1 504-24-5	89415-87-2 128826 2758-42-1 030819 94-82-6 030801 94-75-7 030001 15165-67-0 031402 120-32-1 062201 504-24-5 069201 30560-19-1 103301 57960-19-7 006329 60-35-5 111101 135410-20-7 099050 34256-82-1 121601 135158-54-2 061402	Not Likely To Be Carcinogenic To Humans  Group DNot Classifiable as to Human Carcinogenicity  Not Likely To Be Carcinogenic To Humans  Group CPossible Human Carcinogen  To Human Carcinogenicity  Group CPossible Human  Carcinogen  Not Likely To Be Carcinogenicity  Group CPossible Human  Carcinogen  Not Likely To Be Carcinogenic To Humans  Group CPossible Human  Carcinogen  Not Likely To Be Carcinogenic To Humans  Group CPossible Human  Carcinogen  Not Likely To Be Carcinogenic To Humans  Group CPossible Human  Carcinogen  Not Likely To Be Carcinogenic To Humans  Mot Likely To Be Carcinogenic To Humans  Not Likely To Be Carcinogenic To Humans	Not Likely To Be Carcinogenic To Humans	Not Likely To Be Carcinogenic   8/28/2000 NR

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group DNot Classifiable as to			
Acrinathrin	101007-06-1	129141	Human Carcinogenicity	7/15/1996	NR	Not Applicable
			Data Are Inadequate For An			
			Assessment Of Human			
Acrolein	107-02-8	000701	Carcinogenic Potential	3/25/2008	NR	
			Not Likely To Be Carcinogenic			
ADBAC	68424-85-1	069105	To Humans	12/8/1999	NR	Not Applicable
			Multiple Descriptors: Likely to be Carcinogenic to Humans			
			(High Doses); Not Likely to be			Tumors at multiple sites (Stomach, Nose & Thyroid) in Long
			Carcinogenic to Humans (Low			Evans rats (M & F); Established a thyroid hormonal mode of
Alachlor	15972-60-8	090501	Doses)	6/27/1997	MOE Approach	action for thyroid tumors in rats.
Audenoi	10072 00 0	030301	Group EEvidence of Non-	0/21/1001	MOL Approach	action for thyroid turnors in ratio.
Aldicarb	116-06-3	098301	carcinogenicity for Humans	7/17/2002	NR	Not Applicable
/ Idiodib	110 00 0	000001	Data Are Inadequate for an	771772002	IVIX	Τιστ προποασίο
			Assessment of Human			
Ametryn	834-12-8	080801	Carcinogenic Potential	9/17/2004	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Amicarbazone	129909-90-6	114004	To Humans	8/10/2005	ND	Not Applicable
Afficarbazone	129909-90-0	114004	Not Likely To Be Carcinogenic	6/10/2003	INIX	Not Applicable
Aminopyralid	150114-71-9	005100	To Humans	7/12/2005	NP	Not Applicable
Aminopyrand	130114-71-3	003100	TOTIUMANS	7712/2003	INIX	Lymphoreticular tumors in CFLP mice (F)
			Suggestive Evidence of			Liver tumors in B6C3F1 mice (F)
Amitraz	33089-61-1	106201	Carcinogenic Potential	7/18/2006	NR	Lung tumors B6C3F1 mice (M)
7 WINGE	00000 01 1	100201	Multiple Descriptors: Not Likely	7710/2000	IVIC	Lang tamore become miles (w)
			To Be Carcinogenic To			
			Humans At Doses That Do Not			Thyroid in Charworth Farms rats (M), Fischer 344 rats (M) &
			Alter Rat Thyroid Hormone			Wistar rats (M & F); Established a thyroid hormonal mode of
Amitrole	61-82-5	004401	Homeostasis	5/11/2006	NR	action for thyroid tumors.
7	0.020	001.01	Not Likely To Be Carcinogenic	0, 1 1, 2000		
Aquashade	2650-18-2	110301	To Humans	9/27/2005	NR	Not Applicable
,			Group CPossible Human			
Asulam	3337-71-1	106901	Carcinogen	12/6/2001	NR	Thyroid & Adrenal tumors in Sprague-Dawley rats (M)
			Not Likely To Be Carcinogenic			Not Applicable; Established a neuroendocrine disruption mode
Atrazine	1912-24-9	080803	To Humans	12/13/2000	NR	of action for mammary tumors in rats.
Avermectin (see Emamectin			Group EEvidence of Non-			
Benzoate)	65195-55-3	122804	carcinogenicity for humans	6/27/1996	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
				DATE	METHOD	
			Data Are Inadequate for an			
A	00040 00 0	440040	Assessment of Human	40/40/4000	ND	Not A - Post to
Azafenidin	68049-83-2	119016	Carcinogenic Potential	10/18/1999	NK	Not Applicable
A	00.50.0	050004	Not Likely To Be Carcinogenic	40/7/4000	ND	Not A - Post I
Azinphos-methyl	86-50-0	058001	To Humans	12/7/1993	NK	Not Applicable
A	404000 00 0	400040	Not Likely To Be Carcinogenic	4/44/4007	ND	Not A - Post I
Azoxystrobin	131860-33-8	128810	To Humans	1/14/1997	NK	Not Applicable
D P I	00704 00 0	405004	Group EEvidence of Non-	40/40/4007	ND	Not A - Post to
Bendiocarb	22781-23-3	105201	carcinogenicity for Humans	12/16/1997	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
D. d	4004 40 4	004004	Sufficient to Assess Human	40/07/0004	ND	
Benfluralin	1861-40-1	084301	Carcinogenic Potential	12/27/2001	NR	Liver tumors in B6C3F1 mice (F)
D	47004.05.0	000404	Group CPossible Human	0/04/0000	04* 000 5 0 (0/4)	Liver tumors in CD-1 mice (M &F)
Benomyl	17804-35-2	099101	Carcinogen	9/21/2000	Q1* = 2.39 E-3 (3/4)	Liver tumors in Swiss SPF mice (M & F)
Daniel Karamanti I	00055 00 0	400000	Not Likely To Be Carcinogenic	44/00/4007	ND	Not A - Post I
Bensulfuron methyl	83055-99-6	128820	To Humans	11/20/1997	NK	Not Applicable
5 "	744 50 0	000004	Not Likely To Be Carcinogenic	0/40/4000	LID.	 
Bensulide	741-58-2	009801	To Humans	6/10/1999	NK	Not Applicable
Dantana	05057.00.0	075000	Group EEvidence of Non-	4/44/4000	ND	Not Applicable
Bentazon	25057-89-0	275200	carcinogenicity for Humans	1/14/1992	NK	Not Applicable
			Libebote he Consideration to			Liver tumors in B6C3F1 Mice (M &F)
Doubling alice the income and	477400 00 7	000070	Likely to be Carcinogenic to	40/40/0005	04* 00705 5 0 (0/4)	Thyroid tumors in B6C3F1 Mice (M)
Benthiavalicarb-isopropyl	177406-68-7	098379	Humans	10/18/2005	$Q1^{\circ} = 6.2795 \text{ E-2} (3/4)$	Uterine tumors in Fisher 344 Rat (F)
Danner Cultaria Asid	00504.00.5	100110	Not Likely To Be Carcinogenic To Humans	7/40/0000	ND	Not Applicable
Benzene Sulfonic Acid	68584-22-5	190116		7/19/2006	NK	Not Applicable
Dennyd Denneste	100 51 4	000504	Not Likely To Be Carcinogenic To Humans		ND	Not Applicable
Benzyl Benzoate	120-51-4	009501		6/28/2007	INK	Not Applicable
Diference	440077 44 0	000500	Not Likely To Be Carcinogenic	0/00/0004	ND	Not Applicable
Bifenazate	149877-41-8	000586	To Humans	8/28/2001	NK	Not Applicable
Difanthuin	00057.04.0	100005	Group CPossible Human	0/40/0000	DfD Annyanah	Urinary bladder & Liver tumors (M) and Lung tumors (F) in Swiss Webster mice
Bifenthrin	82657-04-3	128825	Carcinogen	2/19/2003	RfD Approach	Swiss Medister Wice
			Suggestive Evidence of			
			Carcinogenicity, but Not			
Die ellette vie	504.70.0	004000	Sufficient to Assess Human	40/00/0000	ND	Kida ay tura ay in Consanua Bayday Od OB OB (BB) ay ta (M)
Bioallethrin	584-79-2	004003	Carcinogenic Potential	10/29/2003	NK	Kidney tumors in Sprague-DawleyCrl-CD-SD (BR) rats (M)

CHEMICAL	CAS NO.	PC CODE		_	QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
		1		DATE	METHOD	
			Not Likely To Be Carcinogenic			
Bispyrabac Sodium	125401-92-5	078906	To Humans	8/2/2001	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Bitertanol	55179-31-2	117801	To Humans	11/30/2005	NR	Not Applicable
			Group EEvidence Of Non-			
Borax	1303-96-4	011102	Carcinogenicity For Humans	11/24/1993	NR	Not Applicable
			Group EEvidence Of Non-			
Boric acid	10043-35-3	011001	Carcinogenicity For Humans	11/24/1993	NR	Not Applicable
			Group EEvidence of Non-			
Boron	7440-42-8	128945	carcinogenicity for humans	11/24/1993	NR	Not Applicable
			Group EEvidence Of Non-			
Boron Sodium Oxide	12008-41-2	011107	Carcinogenicity For Humans	6/26/2006	NR	Not Applicable
Boron Sodium Oxide,			Group EEvidence Of Non-			
Tetrahydrate	12280-03-4	011103	Carcinogenicity For Humans	11/24/1993	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Boscolid	188425-85-6	128008	Carcinogenic Potential	11/14/2002	NR .	Thyroid tumors in Wistar rats (M & F)
			Group CPossible Human			Liver tumors in CD-1 mice (M)
Bromacil	314-40-9	012301	Carcinogen	1/13/1993	RfD Approach	Thyroid tumors in Crl:CD (BR) rats (M)
			Group CPossible Human			
Bromoxynil	1689-84-5	035301	Carcinogen	6/24/1998	Q1* = 1.03 E-1 (3/4)	Liver tumors in CD-1 mice (M & F)
			Group CPossible Human			
Bromoxynil octanoate	1689-99-2	035302	Carcinogen	6/24/1998	Q1* = 1.03 E-1 (3/4)	Liver tumors in CD-1 mice (M & F)
			Group EEvidence of Non-			
Bromuconazole	116255-48-2	120503	carcinogenicity for humans	4/24/1995	NR	Not Applicable
			Group EEvidence of Non-			
Bronopol	52-51-7	216400	carcinogenicity for humans	6/12/1995	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Buprofezin	69327-76-0	275100	Carcinogenic Potential	3/15/2000	NR	Liver tumors in CD-1 mice (F)
			Likely to be Carcinogenic to			Tumors at multiple sites: Stomach (F) and Kidney, Nose,
Butachlor	23184-66-9	112301	Humans	2/24/1999	NR	Thyroid (M & F) in Sprague-Dawley rats
			Not Likely To Be Carcinogenic			
Butafenacil	134605-64-4	122004	To Humans	7/11/2003	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION		QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group EEvidence of Non-			
Butylate	2008-41-5	041405	carcinogenicity for humans	11/25/1992	NR	Not Applicable
Cacodylic acid	75-60-5	012501	Group BProbable Human Carcinogen	12/14/1999	Q1* = 6.23 E-2 (3/4)	Urinary bladder tumors in Fischer 344 rats (M & F) Fibrosarcomas in multiple organs in B6C3F1 mice (F)
Cadusafos	95465-99-9	128864	Group EEvidence of Non-carcinogenicity for humans	5/28/1992	NR	Not Applicable
Captafol	2939-80-2	081701	Group BProbable Human Carcinogen	5/19/1987	Q1* = 5.1 E-2 (2/3)	Mammary and Liver tumors in Sprague-Dawley rats (F) Kidney tumors in Sprague-Dawley rats (M & F) Lymphosarcomas & Hemangiosarcomas in CD-1 mice (M & F) Harderian gland tumors in CD-1 mice (M)
Captan	133-06-2	081301	Multiple Descriptors: Likely at prolonged, high-level exposures, but not likely at dose levels that do not cause cytotoxicity and regenerative cell hyperplasia	9/22/2004	NR	Intestinal tumors in CD-1 mice (M & F); Established a cytotoxic and regenerative proliferation mode of action for intestinal tumors.
Carbaryl	63-25-2	056801	Likely to be Carcinogenic to Humans	2/12/2002	Q1* = 8.75 E-4 (3/4)	Vascular tumors in CRL:CD-1 (ICR)BR mice (M)
Carbendazim (MBC)	10605-21-7	128872	Group CPossible Human Carcinogen	4/7/1989	Q1* = 2.39 E-3 (3/4)	Liver tumors in CD-1 mice (M & F) Liver tumors in Swiss SPF (M & F)
Carbofuran	1563-66-2	090601	Not Likely To Be Carcinogenic To Humans	6/17/1997	NR	Not Applicable
Carboxin	5234-68-4	090201	Not Likely To Be Carcinogenic To Humans	6/5/2003	NR	Not Applicable
Carfentrazone-ethyl	128639-02-1	128712	Not Likely To Be Carcinogenic To Humans	5/16/2001	NR	Not Applicable
Chlorantraniliprole	500008-45-7	090100	Not Likely To Be Carcinogenic To Humans	3/4/2009	NR	Not Applicable
Chlordimeform	6164-98-3	059701	Group BProbable Human Carcinogen		Q1* = 1.29 E-1 (3/4)	Hemangioendothelomas in Tif:MAG:SPF mice (M & F)
Chlorethoxyfos	54593-83-8	129006	Group DNot Classifiable as to Human Carcinogenicity	3/9/1995	NR	Not Applicable
Chlorfenapyr	122453-73-0	129093	Suggestive Evidence of Carcinogenicity, but Not Sufficient to Assess Human Carcinogenic Potential	3/18/2003	NR	Tumors at multiple sites (Liver, Histiocytic sarcomas and Testes in M; Uterus in F) in Sprague Dawley rats

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION	REPORT DATE	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Not Likely To Be Carcinogenic			
Chlorflurenol Methyl Ester	2536-31-4	098801	To Humans	7/10/2006	NR	Not Applicable
Chlorimuron-ethyl	90982-32-4	128901	Not Likely To Be Carcinogenic To Humans	2/5/2009	NR	Not Applicable
Chlormequat chloride	999-81-5	018101	Not Likely To Be Carcinogenic To Humans	6/12/2007	NR	Not Applicable
Chloroaniline, p-	106-47-8	017203	Group BProbable Human Carcinogen	4/27/1995	5 Q1* = 1.12 E-1 (3/4)	Spleen tumors in F344/N rats (M) Adrenal tumors in F344/N rats (M & F) Liver tumors in B6C3F1 mice (M) Spleen tumors in B6C3F1 mice (M)
Chloroneb	2675-77-6	027301	Data Are Inadequate for an Assessment of Human Carcinogenic Potential	12/18/2003	NR	Not Applicable
Chlorothalonil	1897-45-6	081901	Likely To Be Carcinogenic To	10/20/1997	MOE Approach	Kidney tumors in CD-1 mice (M), Kidney tumors in Fischer 344 rats (M & F) Kidney tumors in Osborne-Mendel rats (M & F) Forestomach tumors in Fischer 344 rats (M & F) Forestomach tumorsCD-1 mice (M & F)
Chlorpropham	101-21-3	018301	Group EEvidence of Non-carcinogenicity for humans	10/11/1994		Not Applicable
Chlorpyrifos	2921-88-2	059101	Group EEvidence of Non-carcinogenicity for humans	11/23/1993	NR	Not Applicable
Chlorpyrifos methyl	1351032	059102	Not Likely To Be Carcinogenic To Humans	5/17/1999	NR	Not Applicable
Chlorsulfuron	64902-72-3	118601	Group EEvidence of Non- carcinogenicity for humans	7/17/2002	NR	Not Applicable
Chlorthal-dimethyl (DCPA)	1861-32-1	078701	Group CPossible Human Carcinogen	2/10/1995	5 Q1* = 1.49 E-3 (3/4)	Thyroid tumors in Sprague-Dawley rats(M & F) Liver tumors in Sprague-Dawley rats (F) Liver tumors CD-1 mice (F)
Chromic acid	7738-94-5, 18540-29-9	021101	Likely To Be Carcinogenic To Humans	7/1/2009	Q1* = 7.91 x 10-1	Oral Mucosa and Tongue Mouse B6C3F1 (M & F) Small Intestines Rat F344 (M & F); Mutagenesis
Clethodim	99129-21-2	121011	Not Likely To Be Carcinogenic To Humans	9/28/2007	NR	Not Applicable
Clodinafop-propargyl	105512-06-9	125203	Suggestive Evidence of Carcinogenic Potential	2/8/2006	NR	Prostate gland tumors in Tif: RAIf (SPF) rat (M) Liver tumors in Tif:MAGf (SPF) mouse (M &F); Established a PPARa mode of action for liver tumors.

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group CPossible Human			
Clofencet (MON 21200)	82697-71-0	128726	Carcinogen	7/23/1996	RfD Approach	Histiocytic sarcomas in CD-1 mice (F)
			Group CPossible Human			
Clofentezine	74115-24-5	125501	Carcinogen	4/3/1990	Q1* = 3.76 E -2 (3/4)	Thyroid tumors in Sprague-Dawley rats (M)
			Not Likely To Be Carcinogenic			
Clomazone	81777-89-1	125401	To Humans	1/31/2001	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Clopyralid	1702-17-6	117403	To Humans	12/20/1999	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Cloquintocet-mexyl	99607-70-2	700099	To Humans	8/31/1999	NR	Not Applicable
			Group EEvidence of Non-			
Cloransulam-methyl	147150-35-4	129116	carcinogenicity for humans	9/30/1997	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Clothianidin	210880-92-5	044309	To Humans	1/6/2003	NR	Not Applicable
			Likely to be Carcinogenic to			Liver tumors in B6C3F1 mice (M &F)
Cocamide Diethanolamine (DEA)	68603-42-9	224600	Humans	7/25/2001	Q1* = 4.01 E-1 (3/4)	Kidney tumors in B6C3F1 mice (M)
			Group DNot Classifiable As			
Copper Compounds	20427-59-2	023401	To Human Carcinogenicity	6/13/2006	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Coumaphos	56-72-4	036501	To Humans	6/25/1999	NR	Not Applicable
			Group DNot Classifiable as to			
Cresol, p-Chloro-m-	59-50-7	064206	Human Carcinogenicity	11/28/1995	NR	Not Applicable
·			Group DNot Classifiable as to			
Cryolite	15096-52-3	075101	Human Carcinogenicity	12/22/1995	NR	Not Applicable
			Suggestive Evidence of			
Cumyluron	99485-76-4	027902	Carcinogenic Potential	6/11/2008	NR	Liver tumors in B6C3F1 mice (M &F)
			Group CPossible Human			
Cyanazine	21725-46-2	100101	Carcinogen	7/30/1991	Q1* = 1.01 E-0 (2/3)	Mammary gland tumors in Sprague- Dawely rats (F)
			Not Likely To Be Carcinogenic		` ` ` ` ` ` ` ` ` ` ` ` ` ` ` ` ` ` `	
Cyazofamid	120116-88-3	085651	To Humans	6/3/2009	NR	Not Applicable
	440400 77 5	200004	Not Likely To Be Carcinogenic	1/0/165=		N A II I
Cyclanilide	113136-77-9	026201	To Humans	4/9/1997	NK	Not Applicable
			Not Likely To Be Carcinogenic			
Cycloate	1134-23-2	041301	To Humans	9/25/2003	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Cyfluthrin	68359-37-5	128831	To Humans	5/21/2002	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			•	DATE	METHOD	Not Applicable, Established a DDADs made of action for liver
Cybalafan hutul	122000 05 0	000500	Not Likely To Be Carcinogenic To Humans	12/20/2007	ND	Not Applicable; Established a PPARa mode of action for liver
Cyhalofop butyl	122008-85-9	082583		12/20/2007	INK	tumros
Cuth a lath via	C000E 0E 0	100007	Group DNot Classifiable as to		ND	Not Applicable
Cyhalothrin	68085-85-8	128867	Human Carcinogenicity	9/15/1994	INK	Not Applicable
			Data Are Inadequate for an			
			Assessment of Human			
Cyhexatin	13121-70-5	101601	Carcinogenic Potential	4/7/2005	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Cymoxanil	57966-95-7	129106	To Humans	1/2/2003	NR	Not Applicable
			Group CPossible Human			
Cypermethrin	52315-07-8	109702	Carcinogen	9/27/1988	ND	Lung tumors in Alderly Park SPF Swiss mice (F)
Суреппешш	32313-07-0	109702		9/2//1900	INIX	Lung turnors in Alderry Fark SFT Swiss filice (1)
			Not Likely To Be Carcinogenic			
			To Humans at doses that do			
			not cause a mitogenic			Liver tumors in CD-1 mice (M & F); Established a non-
Cyproconazole	94361-06-5	128993	response in the liver	12/4/2007	NR	genotoxic, mitogenic mode of action for liver tumors.
			Not Likely To Be Carcinogenic			
Cyprodinil	121552-61-2	288202	To Humans	1/14/1998	NR	Not Applicable
						Kidney tumors in Wistar rats (M);
						Urinary bladder tumors in Wistar rats(F)
						Urinary bladder tumors & Histicocytic sarcomas in C57BL/6J
			Not Likely To Be Carcinogenic			mice (F); Established a cytotoxicity and regenerative
Cyprosulfamide	221667-31-8	877400	To Humans	2/29/2008	NR	proliferation mode of action for urinary bladder tumors.
			Group EEvidence of Non-			
Cyromazine	66215-27-8	121301	carcinogenicity for humans	1/6/1995	NR	Not Applicable
			Not Likely To Be Carcinogenic			
d-Allethrin (Pynamin Forte)	584-79-2	004005	To Humans	6/27/2007	NR	Not Applicable
						Tumors at multiple sites (Cecum, Kidneys, Liver, Lung, Nose,
						Pancreas, Uterus, Vascular) in Fischer 344 rats (M & F);
			Group BProbable Human			B6C3F1 mice (M & F) Swiss mice (M & F); C57BL mice (F);
Daminozide	1596-84-5	035101	Carcinogen	7/26/1991	Q1* = 8.7 E-3 (2/3)	CD-1 mice (M & F) and Syrian Golden hamster (M)
			Not Likely To Be Carcinogenic	.,_0,.501		
Dantochlor (BCDMH)	118-52-5	028501	To Humans	8/14/2000	NR	Not Applicable
(= = =)		1	Group DNot Classifiable as to			11 11 1111
Dazomet	533-74-4	035602	Human Carcinogenicity	12/7/1993	NR	Not Applicable
	300 7 1 1	500002	aa caroniogomony	12,17,1000		· · · · · · · · · · · · · · · · · · ·

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
		1	Not Likely To Be Carcinogenic			
DDBSA	27176-87-0	098002	To Humans	7/19/2006	NR	Not Applicable
			Group DNot Classifiable as to			
DEET	134-62-3	080301	Human Carcinogenicity	1/4/1996	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Deltamethrin	52918-63-5	097805	To Humans	9/9/2003	NR	Not Applicable
			Group EEvidence of Non-			
Desmedipham	13684-56-5	104801	carcinogenicity for humans	11/20/1995	NR	Not Applicable
			Not Likely To Be Carcinogenic			Not Applicable.; Established a neuroendocrine disruption mode
Diaminochlrotrizine (DACT)	3397-62-4	600158	To Humans	4/5/2002	NR	of action for mammary tumors in rats.
			Not Likely To Be Carcinogenic			
Diazinon	333-41-5	057801	To Humans	6/17/1997	NR	Not Applicable
			Group DNot Classifiable as to			
Dicamba	1918-00-9	029801	Human Carcinogenicity	7/29/1996	NR	Not Applicable
			Group CPossible Human			Liver tumors in Fischer 344 rats (M &F)
Dichlobenil	1194-65-6	027401	Carcinogen	7/18/1995	RfD Approach	Liver tumors in Syrian Golden hamsters (M)
			Not Likely To Be Carcinogenic			
Dichlormid	37764-25-3	900497	To Humans	11/15/2005	NR	Not Applicable
			Group DNot Classifiable as to			
Dichlorobenzamide, 2,6-	2008-88-4	027402	Human Carcinogenicity	11/28/1995	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			Mononuclear cell leukemia in Fisher 344 rats (M)
Dichlorvos	62-73-7	084001	Carcinogenic Potential	3/1/2000	NR	Forestomach tumors in B63F1 mice(F)
						Thyroid (F) and Liver (F & M) & Leydig cell (M) tumors in
			Likely to be Carcinogenic to			Wistar rats
Diclofop-methyl	51338-27-3	110902	Humans	5/24/2000	Q1* = 7.36 E-2 (3/4)	Liver tumors in B6C3F1 mice (M & F)
			Suggestive Evidence Of			
Dicloran	99-30-9	031301	Carcinogenic Potential	5/11/2006	NR	Testes tumors in Wistar Rat (M)
			Not Likely To Be Carcinogenic			
Diclosulam	145701-21-9	129122	To Humans	11/9/1999	NR	Not Applicable
			Group CPossible Human			
Dicofol	115-32-2	010501	Carcinogen	6/24/1992	NR	Liver tumors in B6C3F1 mice (M)
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Dicrotophos	141-66-2	035201	Carcinogenic Potential	10/18/1999	NR	Thyroid tumors in C57BL/10 J CD-1 Alpk mice (M & F)

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
Didecyl dimethyl ammonium			Group EEvidence of Non-	DATE	METHOD	
chloride (DDAC)	7173-51-5	069149	carcinogenicity for Humans	4/11/2000	ND	Not Applicable
chloride (DDAC)	7173-31-3	069149	Not Likely To Be Carcinogenic	4/11/2000	INIX	Not Applicable
Diethanolamine Mefluidide	53780-36-2	114002	To Humans	E/20/2007	ND	Not Applicable
Diethanolamine Menuldide	55760-36-2	114002	Group CPossible Human	5/30/2007	INIX	Not Applicable
Difeneganazala	119446-68-3	120047	Carcinogen	7/27/1004	MOE Approach	Liver tumors in CD-1 mice (M & F)
Difenoconazole	119440-00-3	128847	Group EEvidence of Non-	1/21/1994	MOE Approach	Liver turnors in CD-1 mice (W & F)
Diference out mostly a sulfate	42222 40.0	100101		E/04/4004	ND	Not Applicable
Difenzoquat methyl sulfate	43222-48-6	106401	carcinogenicity for humans Group EEvidence of Non-	5/24/1994	INK	Not Applicable
Diflubenzuron	35367-38-5	108201	carcinogenicity for humans	4/27/1995	ND	Not Applicable
Dilluberizurori	35367-36-5	100201	Not Likely To Be Carcinogenic	4/27/1995	INK	Not Applicable
Diffusion zonur Codiium	109293-98-3	005107	To Humans	10/6/1998	ND	Not Applicable
Diflufenzopyr Sodiium	109293-90-3	005107	Group CPossible Human	10/6/1990	INK	Not Applicable
Dimethenamid	87674-68-8	129051		0/45/4005	DfD Annragah	Liver turners in Care que Develou rate (MA)
Dimethenamid	8/6/4-68-8	129051	Carcinogen	9/15/1995	RfD Approach	Liver tumors in Sprague-Dawley rats (M)
Dim oth an amid D	100545 44 0	120051	Group CPossible Human	0/07/0000	DfD Annragah	Liver turners in Coregue Develourete (M)
Dimethenamid-P	163515-14-8	120051	Carcinogen	8/27/2008	RfD Approach	Liver tumors in Sprague-Dawley rats (M)
Discoult in in	55000 04 7	440004	Group CPossible Human	4/5/4000	ND	Lung to the control of CD 4 ratios (M)
Dimethipin	55290-64-7	118901	Carcinogen	1/5/1990	INK	Lung tumors in CD-1 mice (M)
Dimethento	CO E4 E	025004	Group CPossible Human	2/20/2000	DfD Annragah	Hemolymphoreticular tumors in B6C3F1 mice (M)
Dimethoate	60-51-5	035001	Carcinogen  Not Likely To Be Carcinogenic	3/26/2002	RfD Approach	Spleen, Skin, Lymphnode tumors in Wistar rats (M)
Discrete and and b	440400 70 5	00000	,	F/40/4000	ND	Niet Applicable
Dimethomorph	110488-70-5	268800	To Humans	5/13/1998	NK	Not Applicable
Discrete access	000 00 0	004004	Suggestive Evidence of	40/04/0000	ND	Niet Ameliaakia
Dimethoxane	828-00-2	001001	Carcinogenic Potential	12/21/2000	INK	Not Applicable
Discrete discrete	445 40 0	000000	Group DNot Classifiable as to		ND	Niet Ameliaakia
Dimethyl ether	115-10-6	900382	Human Carcinogenicity	1/12/1994	NK	Not Applicable
Discontinuole etais	40070 00 0	000045	Not Likely To Be Carcinogenic	0/00/0000	ND	Niet Applicable
Dimethylhydantoin	16079-88-2	006315	To Humans	8/28/2000	INK	Not Applicable
<b>D</b>	00000 45 0	000004	Group EEvidence of Non-	0/00/4004	ND	Not A - Post I
Dinocap	39300-45-3	036001	carcinogenicity for Humans	6/22/1994	NK	Not Applicable
D' I	00.05.7	007505	Group CPossible Human	0/40/4000	LID.	Li cot con i OD to i o (E)
Dinoseb	88-85-7	037505	Carcinogen	6/19/1986	NK	Liver tumors in CD-1 mice (F)
D:	105050 75 5	0.1.10.10	Not Likely To Be Carcinogenic	0/5/055		
Dinotefuran	165252-70-0	044312	To Humans	3/5/2004	NK	Not Applicable
	100.00.4	200504	Not Likely To Be Carcinogenic	4/4/465		N A II II.
Diphenylamine	122-39-4	038501	To Humans	4/1/1997	NK	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group EEvidence of Non-			
Diquat dibromide	85-00-7	032201	carcinogenicity for Humans	5/12/1994	NR	Not Applicable
			Likely To Be Carcinogenic To			Oral Mucosa and Tongue Mouse B6C3F1 (M & F)
Disodium Dichromate Dihydrate		068306	Humans	7/1/2009	Q1* = 7.91 x 10-1	Small Intestines Rat F344 (M & F); Mutagenesis
			Not Likely To Be Carcinogenic			
Disodium methanearsonate	144-21-8	013802	To Humans	7/26/2000	NR	Not Applicable
			Group EEvidence of Non-			
Disulfoton	298-04-4	032501	carcinogenicity for Humans	4/21/1997	NR	Not Applicable
			Suggestive Evidence of			
Dithianon	3347-22-6	099201	Carcinogenic Potential	2/23/2006	NR	Kidney tumors in Sprague Dawley rats (F)
			Group EEvidence of Non-			
Dithiopyr (MON 7200)	97886-45-8	128994	carcinogenicity for Humans	5/29/1997	NR	Not Applicable
						Urinary bladder tumors in Wistar rats (M&F)
						Kidney tumors in Wistar rats (M)
Diuron	330-54-1	035505	Known/Likely	5/8/1997	Q1* = 1.91 E-2 (3/4)	Mammary tumors in NMRI mice (F)
			Not Likely To Be Carcinogenic			
Dodine	2439-10-3	044301	To Humans	1/24/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Ecolyst	274671-61-3	069089	To Humans	10/19/1999	NR	Not Applicable
Emamectin Benzoate (Deoxy			Not Likely To Be Carcinogenic			
Avermectin)	137512-74-4	122806	To Humans	3/19/1998	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Endosulfan	115-29-7	079401	To Humans	1/31/2000	NR	Not Applicable
			Not Likely To Be Carcinogenic		<u>_</u>	
Endothall	145-73-3	038901	To Humans	10/23/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Endothall Amine Salt	66330-88-9	038905	To Humans	10/23/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Endothall dipotassium salt	2164-07-0	038904	To Humans	10/23/2008	NR	Not Applicable
						Liver tumors in C57BL/6N CrlBr mice (M & F)
	106325-08-0,		Likely to be Carcinogenic to			Liver and Adrenal (M & F) and ovarian (F) tumors in Wistar
Epoxiconazole	133855-98-8	123909	Humans	1/24/2001	Q1* = 3.04E-2 (3/4)	rats
			Suggestive Evidence of			
			Carcinogenicity, but Not			
Edition 2	00404 00 0	004007	Sufficient to Assess Human	40/0/0000	LID.	16 by 1 may 1 0 may 1 0 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
Esbiothrin	28434-00-6	004007	Carcinogenic Potential	12/2/2003	NK	Kidney tumors in Sprague-Dawley Crl-CD-SD(BR) rats (M)

CHEMICAL	CAS NO.	PC CODE		REPORT DATE	QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group EEvidence of Non-	IDATE	METHOD	
Esfenvalerate	66230-04-4	109303	carcinogenicity for Humans	7/1/1996	ND	Not Applicable
Esterivalerate	00230-04-4	109303	Suggestive Evidence of	7/1/1990	INIX	Not Applicable
Ethaboxam	162650-77-03	000005	Carcinogenic Potential	3/23/2006	ND	Loudig cell tumore in Carague Doulloy rate (M)
Elliaboxalli	102030-77-03	090205		3/23/2000	INK	Leydig cell tumors in Sprague Dawley rats (M)  Mammary, Urinary bladder & Kidney tumors in Fischer 344
Ethalfluralin	55283-68-6	110101	Group CPossible Human	0/4.4/4.00.4	04* 00 5 2 (2/4)	
Emamuraiin	55283-68-6	113101	Carcinogen		Q1* = 8.9 E-2 (3/4)	rats (M & F)
Cth anh an	10070 07 0	000004	Group DNot Classifiable as to		ND	Net Applicable
Ethephon	16672-87-0	099801	Human Carcinogenicity	8/15/1994	INK	Not Applicable
	500 40 0	050404	Group EEvidence of Non-	4/00/400	NB	N . A . B . I .
Ethion	563-12-2	058401	carcinogenicity for humans	1/26/1994	NR	Not Applicable
			Group DNot Classifiable as to			
Ethofumesate	26225-79-6	110601	Human Carcinogenicity	2/24/1994	NR	Not Applicable
			Likely to be Carcinogenic to			Adrenal tumors in Sprague-Dawley rats (M)
Ethoprop	13194-48-4	041101	Humans	10/7/1998	Q1* = 2.81 E-2 (3/4)	Thyroid tumors in Sprague-Dawley & Fischer 344 rats (M)
Ethyl dipropylthiocarbamate			Not Likely To Be Carcinogenic			
(EPTC)	759-94-4	041401	To Humans	8/31/1999	NR	Not Applicable
			Group BProbable Human			Thyroid tumors in Fischer 344 rats (M & F)
Ethylene thiourea (ETU)	96-45-7	600016	Carcinogen	7/7/1999	Q1* = 6.01 E-2 (3/4)	Pituitary and Liver tumors in B6C3F1 mice (M & F)
			Multiple Descriptors: Not Likely			Thyroid tumors in Sprague-Dawley rats (M & F); Established a
Etofenprox	80844-07-1	128965	Below a Defined Dose Range	2/8/2006	NR	thyroid hormone disruption mode of action for thyroid tumors.
			Not Likely To Be Carcinogenic			
Etoxazole	153233-91-1	107091	To Humans	8/7/2003	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Famoxadone	131807-57-3	113202	To Humans	4/16/2003	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Fenamidone	161326-34-7	046679	To Humans	7/12/2002	NR	Not Applicable
			Group EEvidence of Non-			1
Fenamiphos	22224-92-6	100601		11/23/1993	NR	Not Applicable
Fenarimol	60168-88-9	206600		9/5/2001	NR	Not Applicable
	32.122.22		1	2, 2, 200		11. 11
Fenazaguin	120928-09-8	044501		5/15/2007	NR	Not Applicable
	120020 00 0	- 11001		5, 15,2001		
Fenhuconazole	114369-43-6	129011	1	4/15/1996	01* = 3 50 F-3 (3/4)	
Fenamiphos  Fenarimol  Fenazaquin  Fenbuconazole	22224-92-6 60168-88-9 120928-09-8 114369-43-6	100601 206600 044501 129011	Group EEvidence of Non- carcinogenicity for Humans Not Likely To Be Carcinogenic To Humans Not Likely To Be Carcinogenic To Humans Group CPossible Human Carcinogen	11/23/1993 9/5/2001 5/15/2007 4/15/1996	NR	Not Applicable  Not Applicable  Not Applicable  Thyroid tumors in Sprague-Dawley rats (M)  Liver tumors in CD-1 mice (M & F)

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION		QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group EEvidence of Non-			
Fenbutatin-oxide	13356-08-6	104601	carcinogenicity for Humans	3/2/1993	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Fenhexamide	126833-17-8	090209	To Humans	3/4/1999	NR	Not Applicable
Fenitrothion	122-14-5	105901	Group EEvidence of Non-carcinogenicity for Humans	7/13/1993	NR	Not Applicable
Fenoxycarb	72490-01-8	125301	Likely to be Carcinogenic to Humans	4/1/1996	Q1* = 7.00 E-2 (3/4)	Lung tumors & Harderian gland tumors in CD-1 mice (M)
			Not Likely To Be Carcinogenic			
Fenpropathrin	39515-41-8	127901	To Humans	12/22/2003	NR	Not Applicable
			Suggestive Evidence Of			
Fenpropidin	67306-00-7	012305	Carcinogenic Potential	6/9/2009	NR	Pancreas Rat Sprague-Dawley (M); No
			Not Likely To Be Carcinogenic			
Fenpropimorph	67564-91-4	121402	To Humans	10/19/2005	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Fenpyroximate	134098-61-6	129131	To Humans	2/19/1997	NR	Not Applicable
			Group EEvidence of Non-			
Fenthion	55-38-9	053301	carcinogenicity for Humans	3/11/1996	NR	Not Applicable
			Group EEvidence of Non-			
Fenvalerate	51630-58-1	109301	carcinogenicity for Humans	2/10/2003	NR	Not Applicable
						Thyroid tumors & Hemangiomas in Fischer 344 rats &
			Likely to be Carcinogenic to			CD rats (M)
Ferbam	128-04-1	034801	Humans	4/6/2000	NR	Lung tumors in B6C3F1 mice (F)
			Group CPossible Human			
Fipronil	120068-37-3	129121	Carcinogen	7/18/1995	RfD Approach	Thyroid tumors in CD rats (M & F)
			Not Likely To Be Carcinogenic		l	
Flazasulfuron	104040-78-0	119011	To Humans	11/16/2005	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			Nasolacrimal duct tumors in Wistar rats (F)
			Sufficient to Assess Human			Lung tumors in CD-1 mice (M & F); Established a mitogenic
Flonicamid	158062-67-0	128016	Carcinogenic Potential	2/24/2005	NR	mode of action for mouse lung tumors.
			Not Likely To Be Carcinogenic	_ /2 / /2 2 2 -		
Florasulam	145701-23-1	129108	To Humans	5/31/2007	NR	Not Applicable
		400005	Not Likely To Be Carcinogenic	0/4.4/055		N A II I.
Fluazifop	69806-50-4	122805	To Humans	6/14/2004	NK	Not Applicable
Flora-ifan D. Dodol	70044 40 0	400000	Not Likely To Be Carcinogenic	0/40/0000	ND	Net Applicable
Fluazifop-P-Butyl	79241-46-6	122809	To Humans	9/19/2008	NK	Not Applicable

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			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			Thyroid tumors in Sprague-Dawley (M)
Fluazinam	79622-59-6	129098	Carcinogenic Potential	3/29/2001	NR	Liver tumors in CD-1 mice (M)
			Not Likely To Be Carcinogenic			
Flubendiamide		027602	To Humans	4/3/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Flucarbazone-sodium	181274-17-9	114009	To Humans	7/19/2000	NR	Not Applicable
			Group DNot Classifiable as to			
Fludioxonil	131341-86-1	071503	Human Carcinogenicity	9/19/1996	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Flufenacet (Thiaflumide)	142459-58-3	121903	To Humans	7/16/1997	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Flufenoxuron	101463-69-8	108203	To Humans	8/15/2006	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Flufenpyr-ethyl	188489-07-8	108853	To Humans	6/8/2003	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Flumetralin	62924-70-3	123001	To Humans	6/21/2007	NR	Not Applicable
			Group EEvidence of Non-			
Flumetsulam (XRD-498)	98967-40-9	129016	carcinogenicity for Humans	3/24/1993	NR	Not Applicable
			Group EEvidence of Non-			
Flumiclorac pentyl	87546-18-7	128724	carcinogenicity for Humans	9/7/1994	NR	Not Applicable
	103361-09-7,		Not Likely To Be Carcinogenic			
Flumioxazin	141490-50-8	129034	To Humans	2/22/2001	NR	Not Applicable
			Group CPossible Human			
Fluometuron	2164-17-2	035503	Carcinogen	8/28/1996	Q1* = 1.80 E-2 (3/4)	Lung tumors (M) & Lymphocytic lymphomas (F) in CD-1 mice
			Not Likely To Be Carcinogenic			Liver tumors in C57Bl/6 mice (M & F); Established a mitogenic
Fluopicolide	239110-15-7	027412	To Humans	12/12/2006	RfD Approach	mode of action for liver tumors in mice.
			Not Likely To Be Carcinogenic			
Fluoxastrobin	361377-29-9	028869	To Humans	1/24/2005	NR	Not Applicable
			Group EEvidence of Non-			
Fluridone	59756-60-4	112900	carcinogenicity for Humans	7/1/1985	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Fluroxypyr	81406-37-3	128968	To Humans	6/26/2003	NR	Not Applicable
Fluroxypyr acid (see also PC			Not Likely To Be Carcinogenic			
Code 128968)	69377-81-7	128959	To Humans	6/26/2003	NR	Not Applicable

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			Not Likely To Be Carcinogenic			
Flurprimidol	56425-91-3	125701	To Humans	9/29/2005	NR	Not Applicable
			Likely to be Carcinogenic to			Pancreatic tumors in Sprague-Dawley rats (M)
Fluthiacet methyl	117337-19-6	108803	Humans	11/20/1998	Q1* = 2.07 E-1 (3/4)	Liver tumors in CD-1 mice (M & F)
			Group EEvidence of Non-			
Flutolanil	66332-96-5	128975	carcinogenicity for Humans	6/9/1994	NR	Not Applicable
· ratera	00002 00 0		Not Likely To Be Carcinogenic	0,0,100		Trott pp nousio
Flutriafol	76674-21-0	128940	To Humans	6/1/2009	NR	Not Applicable
· ratifact			. o mamano	0, 1,2000		Duodenum tumors in CD-1 mice (M & F) and
			Group BProbable Human			B6C3F1 mice (M & F)
Folpet	133-07-3	081601	Carcinogen	8/19/2003	Q1* = 1.86 E-3 (3/4)	Skin tumors in B6C3F1 mice (M)
			Not Likely To Be Carcinogenic			Liver tumors in CD-1 mice (M & F); Established a PPARa
Fomesafen	108731-70-0	123802	To Humans	11/3/2005	NR	mode of action for liver tumors
			Group EEvidence of Non-			
Fonofos	944-22-9	041701	carcinogenicity for Humans	11/10/1993	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Forchlorfenuron	68157-60-8	128819	To Humans	3/11/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Formasulfuron	173159-57-4	122020	To Humans	9/19/2001	NR	Not Applicable
			Group EEvidence of Non-			
Formetanate hydrochloride	23422-53-9	097301	carcinogenicity for Humans	5/20/1996	NR	Not Applicable
·			Not Likely To Be Carcinogenic			
Fosetyl-Al	39148-24-8	123301	To Humans	4/22/1999	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Fosthiazate	98886-44-3	129022	To Humans	9/15/2003	NR	Not Applicable
			Likely to be Carcinogenic to			Tumors at multiple sites (Liver, Lung, Stomach, Testes) in
Furiazole (MON 13900)	121776-33-8	911596	Humans	10/15/1999	Q1* = 2.74 E-2 (3/4)	Sprague-Dawley rats (M&F) & CD-1 mice (M & F)
			Group BProbable Human			
Furmecyclox	60568-05-0	122601	Carcinogen	7/3/1985	Q1* = 2.98 E-2 (2/3)	Liver & Urothelial tumors in Sprague-Dawley rats (M & F)
			Not Likely To Be Carcinogenic			
Gamma Cyhalothrin	76703-62-3	128807	To Humans	3/1/2004	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Glufosinate-ammonium	77182-82-2	128850	To Humans	5/17/1999	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Glutaraldehyde	111-30-8	043901	To Humans	5/18/2006	NR	Not Applicable
			Group EEvidence of Non-			
Glyphosate	1071-83-6	417300	carcinogenicity for Humans	12/31/1991	NR	Not Applicable

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			Not Likely To Be Carcinogenic			
Halosulfuron methyl (MON 1200)	100784-20-1	128721	To Humans	2/26/1998	NR	Not Applicable
			Group BProbable Human			
Haloxyfop-methyl	690806-40-2	125201	Carcinogen	9/18/1989	Q1* = 7.39 E+0 (2/3)	Liver tumors in B6C3F1 mice (M & F)
			Group CPossible Human			
Hexaconazole	79983-71-4	128925	Carcinogen	1/21/1999	Q1* = 1.6 E-2 (3/4)	Leydig cell tumors in Wistar (Alpk:APfSD) rats (M)
		021101;	Likely to be Carcinogenic to			Oral mucosa & Tongue tumors in F344 rats (M & F) Intestinal (duodenum, jejunum, and ileum) tumors in B6C3F1
Hexavalent Chromium (CrVI)	18540-29-9	068302	Humans		Q1* = 7.91 E-1 (3/4)	mice (M & F)
			Group DNot Classifiable as to			
Hexazinone	51235-04-2	107201	Human Carcinogenicity	7/27/1994	NR	Not Applicable
			Likely To Be Carcinogenic To			Liver tumors in B6C3F1 mice (F) Mammary Gland tumors
Hexythiazox	78587-05-0	128849	Humans	8/12/2009	RfD Approach	(fibroadenomas) in Fisher 344 Rats (M); Not Applicable
			Not Likely To Be Carcinogenic			
HOE107892	135590-91-9	811800	To Humans	11/24/1998	NR	Not Applicable
			Group CPossible Human			
Hydramethylnon	67485-29-4	118401	Carcinogen	3/28/1991	RfD Approach	Lung tumors in CD-1 mice (F)
			Group CPossible Human			
Hydrogen cyanamide	420-04-2	014002	Carcinogen		Q1* = 6.64 E-2 (3/4)	Ovarian tumors in CRL:CD-1 (ICR)BR mice (F)
			Group DNot Classifiable as to			
Hydroprene	41096-46-2	486300	Human Carcinogenicity	6/8/1995	NR	Not Applicable
			Likely to be Carcinogenic to			Liver & Thyroid tumors in Wistar rats (M)
Imazalil	35554-44-0	111901	Humans		Q1* = 6.11 E-2 (3/4)	Liver tumors in Swiss albino mice (M)
			Group DNot Classifiable as to			
Imazamethabenz	81405-85-8	128842	Human Carcinogenicity	6/11/1987	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Imazamox	114311-32-9	129171	To Humans	2/27/1997	NR	Not Applicable
			Group EEvidence of Non-			
Imazapic	81334-60-3	129041	carcinogenicity for Humans	9/27/1995	NR	Not Applicable
			Group EEvidence of Non-			
Imazapyr	81334-34-1	128821	carcinogenicity for Humans	10/5/1995	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Imazaquin Acid	81335-37-7	128848	To Humans	10/31/2005	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Imazaquin ammonium	81335-47-9	128840	To Humans	10/31/2005	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Imazaquin Sodium	81335-46-8	129023	To Humans	10/31/2005	NR	Not Applicable

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			Not Likely To Be Carcinogenic			
Imazethapyr	81335-77-5	128922	To Humans	1/31/2002	NR	Not Applicable
			Group EEvidence of Non-			
Imidacloprid	105827-78-9	129099	carcinogenicity for Humans	11/10/1993	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Indoxacarb	173584-44-6	067710	To Humans	7/17/2000	NR	Not Applicable
mackacais	170007 44 0	007710	Multiple Descriptors: Not Likely		THE	Not Applicable
			to be Carcinogenic to Humans			Thyroid tumors in Fischer 344 rats (M)
			at doses that do not alter rat			Thyroid tumors in B6C3F1 mice (M); Established a thyroid
lodomethane	74-88-4	000011	thyroid hormone homeostasis	11/10/2005	RfD Approach	hormonal mode of action for thyroid tumors.
Todol Hot Harie	7 7 00 7	000011	Not Likely To Be Carcinogenic	11/10/2000	тав трргоцоп	normal mode of dealers for anytoid turners.
lodosulfuran	144550-36-7	122021	To Humans	1/5/2004	NR	Not Applicable
			Not Likely To Be Carcinogenic	1,0,00		The state of the s
Ipoconazole	125225-28-7	125618	To Humans	5/28/2008	3	
			Likely to be Carcinogenic to			Liver (M & F) & Ovarian luteomas (F) in CD-1 mice
Iprodione	36734-19-7	109801	Humans	2/26/1998	Q1* = 4.39 E-2 (3/4)	Leydig cell tumors in Crl:CD(SD)BR rats (M)
			Likely to be Carcinogenic to			Tumors at multiple sites (Osteosarcomas, Urinary bladder,
Iprovalicarb	140923-17-7	098359	Humans	4/11/2002	Q1* = 4.47E-4 (3/4)	Uterus, Thyroid) in Wistar (Hsd/WIN:WU) rats (M & F)
	110000		Group EEvidence of Non-			Constant Con
Isofenphos	25311-71-1	109401	carcinogenicity for Humans	1/13/1998	NR	Not Applicable
·			Group CPossible Human			
Isophorone	78-59-1	047401	Carcinogen	9/2/1999	Q1* = 6.08 E-4 (3/4)	Preputial gland tumors in F344/N rats (M)
			Suggestive Evidence of			
Isoxaben	82558-50-7	125851	Carcinogenic Potential	10/7/2008	NR	Liver tumors in B6C3F1 mice (M & F)
			Not Likely To Be Carcinogenic			
Isoxadifen-ethyl	163520-33-0	823000	To Humans	1/29/2001	NR	Not Applicable
						Liver (M & F) & Thyroid (M) tumors in CrL:CD(SD)
			Likely to be Carcinogenic to			BR VAF/Plus rats
Isoxaflutole	141112-29-0	123000	Humans	8/6/1997	Q1* = 1.02 E-2 (3/4)	Liver tumors in CD-1 mice (M & F)
			Not Likely To Be Carcinogenic			
Kasugamycin	6980-18-3	230001	To Humans	8/17/2005	NR	Not Applicable
			Group DNot Classifiable as to			
Kathon 886	55965-84-9	107106	Human Carcinogenicity		MOE Approach	Not Applicable
			Not Likely To Be Carcinogenic			
KBR 3023	119515-38-7	070705	To Humans	6/9/1999	NR	Not Applicable

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			Likely to be Carcinogenic to			
Kresoxim-methyl	143390-89-0	129111	Humans	8/19/1999	Q1* = 2.90 E-3 (3/4)	Liver tumors in Wistar rats (M & F)
			Multiple Descriptors: Likely to		,	· · ·
			be Carcinogenic in Humans at			
			High Doses. Not Likely to be			Liver neoplastic nodules in Sprague-Dawley rats (M & F)
			Carcinogenic to Humans at			Liver tumors in CD-1 mice (M &F); Established a PPARa mode
Lactofen	77501-63-4	128888	Low Doses	10/17/2006	MOE approach	of action for liver tumors
			Group DNot classifiable as to			
Lambda cyhalothrin	91465-08-6	128897	Human Carcinogenicity	9/12/2002	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Lindane	58-89-9	009001	Carcinogenic Potential	11/29/2001	NR	Lung tumors in CD-1, Pseudoagouti, & Agouti mice (F)
			Group CPossible Human			Testicular tumors in CD rats (M)
Linuron	330-55-2	035506	Carcinogen	11/20/2001	NR	Liver tumors in CD-1 mice (M & F)
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			Liver, Oral palate & Nosetumors in Fischer 344 rats (M & F)
Malathion	121-75-5	057701	Carcinogenic Potential	4/28/2000	NR	Liver tumors in B6C3F1 mice (M & F)
			Group EEvidence of Non-			
Maleic hydrazide	123-33-1	051501	carcinogenicity for Humans	11/10/1993		Not Applicable
			Group BProbable Human		Q1* = 6.01 E-2 (3/4)	
Mancozeb	8018-01-7	014504	Carcinogen	7/7/1999	Based on ETU	Thyroid tumors in Crl:CD(BR) rats (M & F)
			Not Likely To Be Carcinogenic			
Mandipropamid	374726-62-2	036602	To Humans	1/21/2009		Not Applicable
			Group BProbable Human		Q1* = 6.01 E-2 (3/4)	Liver tumors in B6C3F1 mice ( M & F) No acceptable study in
Maneb	12427-38-2	014505	Carcinogen	7/7/1999	Based on ETU	rats
MB46513 (photodegradate of			Not Likely To Be Carcinogenic			
Fipronil)	120067-83-6	600050	To Humans	12/6/2000	NR	Not Applicable
			Not Likely To Be Carcinogenic			
MCPA + Salts	94-74-6	030501	To Humans	10/29/2003	NR	Not Applicable
			Not Likely To Be Carcinogenic		l	
MCPB Acid	94-81-5	019201	To Humans	10/1/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
MCPB Sodium Salt	6062-26-6	019202	To Humans	10/1/2008	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE		_	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Mecoprop-P	16484-77-8	129046	Carcinogenic Potential	3/13/2003	NR	Liver tumors in B6C3F1/CrlBR mice (F)
			Not Likely To Be Carcinogenic			
Mefenoxam	70630-17-0	113502	To Humans	5/17/2000	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Mefluidide	53780-34-0	114001	To Humans	5/30/2007	NR	Not Applicable
			Group DNot Classifiable as to			
Melamine	108-78-1	777201	Human Carcinogenicity	7/21/1993	NR	Not Applicable
			Likely to be Carcinogenic to			Liver tumors in Fisher 344 rats (F)
Mepanipyrim	110235-97-7	288203	Humans	4/20/2004	Q1* = 1.35 E-2 (3/4)	Liver tumors in B6C3F1 mice (M & F)
			Not Likely To Be Carcinogenic			
Mepiquat Chloride	24307-26-4	109101	To Humans	2/19/2003	NR	Not Applicable
			Group CPossible Human			
Mercaptobenzothiazole, 2-	149-30-4	051701	Carcinogen	11/19/1992	RfD Approach	Adrenal (M & F) and Pituitary (F) tumors in F344/N rats
			Not Likely To Be Carcinogenic			
Mesosulfuron methyl	208465-21-8	122009	To Humans	3/4/2004	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Mesotrione	104206-82-8	122990	To Humans	4/12/2001	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Metaflumizone	139968-49-3	281250	To Humans	1/24/2006	NR	Not Applicable
			Group EEvidence of Non-			
Metalaxyl	57837-19-1	113501	carcinogenicity for Humans	4/20/1994	NR	Not Applicable
			Suggestive Evidence of			Liver tumors in Sprague Dawley rats (F)
Metaldehyde	108-62-3	053001	Carcinogenic Potential	6/23/2005	NR	Liver tumors in CD-1 mice (M & F)
			Likely to be Carcinogenic to			Hemangiosarcomas in Hsd/Ola: Wistar rats (M)
Metam Potassium	137-41-7	039002	Humans	5/14/2009	Q1* = 1.98 E-1(3/4)	Angiosarcomas in C57BL/10JfCD-1/Alpk mice (M & F)
			Likely to be Carcinogenic to			Hemangiosarcomas in Hsd/Ola: Wistar rats (M)
Metam sodium	137-42-8	039003	Humans	5/14/2009	Q1* = 1.98 E-1(3/4)	Angiosarcomas in C57BL/10JfCD-1/Alpk mice (M & F)
			Not Likely To Be Carcinogenic			Liver tumors in CD-1 mice (M & F); Established a mitogenic
Metconazole	125116-23-6	125619	To Humans	4/14/2006	NR	mode of action for liver tumors in mice
			Not Likely To Be Carcinogenic			
Methamidophos	10265-92-6	101201	To Humans	10/6/1997	NR	Not Applicable
			Group CPossible Human			
Methidathion	950-37-8	100301	Carcinogen	2/19/1988	NR	Liver tumors in CD-1 mice (M)

CAS NO.	PC CODE				TUMOR SITES/ STRAIN/ SPECIES/ SEX
				IMETHOD	
2022 65 7	100501			DfD Annragah	Not Applicable
2032-65-7	100501		3/2/1993	RID Approach	Not Applicable
10750 77 5	000004		40/05/4006	ND	Not Applicable
10/52-//-5	090301		10/25/1996	INK	Not Applicable
101050 50 4	101007		7/4/4000	ND	Not Applicable
101000-00-4	121027		7/1/1998	INK	Not Applicable
74 92 0	052201		9/4/1002	ND	Not Applicable
14-03-9	053201		0/4/1992	INK	Not Applicable
556 61 6	069102		4/20/2000	ND	Not Applicable
330-01-0	000103		4/30/2008	INIX	Not Applicable
208 00 0	052501		12/1/1007	ND	Not Applicable
290-00-0	055501		12/1/1997		Not Applicable
0006-42-2	01/1601		7/7/1000	, ,	Thyroid tumors in Crl:CD(BR) rats (M & F)
9000-42-2	014001		1/1/1999	Dased on LTO	Thyroid turnors in Ch.CD(Dix) rats (W & r)
		,			
					Not Applicable; Established a mitogenic mode of action for
240444-70-6	100700	_	7/26/2007	NR	liver tumors in rats.
240444-70-0	109709		1/20/2001	INIX	liver turnors in rats.
51218-45-2	108801	i ·	11/16/1994	MOF Approach	Liver tumors in Charles River CD (SD)BR rats (F)
31210 432	100001		11/10/1334	MOL Approach	Elver turnors in Orianes (tiver ob (ob)bit rats (i)
220899-03-6	000325		7/6/2006	NR	Liver Tumors in CD-1 Mice (M)
220000 00 0	000020			IVIC	Erver Famore in OB 1 who (w)
21087-64-9	101101			NR	Not Applicable
21007 010	101101		G/ 10/ 1000	T T T	Trock / pp///ddb//
74223-64-6	122010		3/14/2002	NR	Not Applicable
7 1220 01 0	122010		0/11/2002		Trock reprised to
7786-34-7	015801		5/17/2000	NR	Not Applicable
1100011	0.000.		0,11,200		Liver tumors in CD-1 mice (M & F)
113-48-4	057001	i ·	6/7/1995	RfD Approach	Thyroid tumors in Crl:CDBR rats (M)
1.12.13.		3	2 300		Tumors at multiple sites (Liver, Kidney, Testes, Uterus) in CD
		Group BProbable Human			rats (M & F)
136-45-8	047201		11/12/2002	Q1* = 1.6 E-3 (3/4)	Liver (M) & Lung (F) tumors in CD-1 mice
.55 .55			.,,_		
77-48-5	006317	To Humans	8/28/2000	NR	Not Applicable
	2032-65-7 16752-77-5 161050-58-4 74-83-9 556-61-6 298-00-0 9006-42-2 240444-70-6 51218-45-2 220899-03-6 21087-64-9 74223-64-6 7786-34-7 113-48-4	2032-65-7 100501 16752-77-5 090301 161050-58-4 121027 74-83-9 053201 556-61-6 068103 298-00-0 053501 9006-42-2 014601 240444-70-6 109709 51218-45-2 108801 220899-03-6 000325 21087-64-9 101101 74223-64-6 122010 7786-34-7 015801 113-48-4 057001	Group DNot Classifiable as to Human Carcinogenicity  Group EEvidence of Non-carcinogenicity for Humans  Not Likely To Be Carcinogenic To Humans  There are insufficient data to characterize the cancer risk of MITC.  Not Likely To Be Carcinogenic To Humans  Group BProbable Human  Group BProbable Human  Group BProbable Human  Carcinogen  Not Likely To Be Carcinogenic To Humans At Doses That Do Not Result In A Mitogenic Response  Group CPossible Human  51218-45-2 108801 Carcinogen  Suggestive Evidence of Carcinogenic Potential  Group DNot Classifiable as to Human Carcinogenic To Humans  Not Likely To Be Carcinogenic To Human Carcinogenic Potential  Group DNot Classifiable as to Human Carcinogenicity  Not Likely To Be Carcinogenic To Humans  Not Likely To Be Carcinogenic To Humans  O15801 To Humans  Not Likely To Be Carcinogenic To Humans  Group CPossible Human  Carcinogen  To Humans  O15801 To Humans  Group CPossible Human  Carcinogen  Group BProbable Human  Carcinogen  Group BProbable Human  Carcinogen  Not Likely To Be Carcinogenic	Carcinogen   Carcinogenic   Carcin	Carcinogenicity   Carcinogenic   C

CHEMICAL	CAS NO.	PC CODE		_	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Suggestive Evidence of Carcinogenicity, but Not Sufficient to Assess Human			
Molinate	2212-67-1	041402	Carcinogenic Potential	12/14/2000	NR	Kidney & Testicular tumors in Crl:CD(SD)BR rats (M)
MON 4660	71526-07-3	600046	Likely to be Carcinogenic to Humans	12/9/1999	Q1* = 4.85 E-2 (3/4)	Liver (M & F), Stomach & Bile duct (M) tumors in Sprague Dawley rats Lung (M) and Liver & Stomach (M & F) tumors in CD-1 mice
Monosodium acid			Not Likely To Be Carcinogenic			
methanearsonate (MMA)	2163-80-6	013803	To Humans	7/26/2000	NR	Not Applicable
MSMA-calcium salt	5902-95-4	013806	Not Likely To Be Carcinogenic To Humans	12/14/2000	NR	Not Applicable
Myclobutanil	88671-89-0	128857	Group EEvidence of Non-carcinogenicity for Humans	6/16/1994	NR	Not Applicable
Naled	300-76-5	034401	Group EEvidence of Non-carcinogenicity for Humans	8/31/1994	NR	Not Applicable
Napropamide	15299-99-7	103001	Not Likely To Be Carcinogenic To Humans	7/7/2005	NR	Not Applicable
Naptalam Sodium Salt	132-67-2	030703	Group DNot Classifiable as to Human Carcinogenicity	9/7/1994	NR	Not Applicable
Nicosulfuron	111991-09-4	129008	Group EEvidence of Non-carcinogenicity for Humans	9/1/1998	NR	Not Applicable
Nitrapyrin	1929-82-4	069203	Likely to be Carcinogenic to Humans	3/26/2005	Q1* = 4.25 E-2 (3/4)	Liver (M & F) & Epididymal (M) tumors in B6C3F mice
Norflurazon	27314-13-2	105801	Group CPossible Human Carcinogen	11/2/1990	NR	Liver tumors in CD-1 mice (M)
Novaluron	116714-46-6	124002	Not Likely To Be Carcinogenic To Humans	2/4/2004		Not Applicable
Orthophenylphenol, Sodium salt (see also PC 064103)	132-27-4	064104	Not Likely To Be Carcinogenic To Humans	10/12/2005		Not Applicable: Not Applicable; Established a cytotoxic mode of action involving oxidative damage to cells and subsequent regenerative hyperplasia for bladder tumors in rats.
Orthophenylphenol, Sodium salt (see also PC 064104)	90-43-7	064103	Not Likely To Be Carcinogenic To Humans	10/12/2005	NR	Not Applicable; Established a cytotoxic mode of action involving oxidative damage to cells and subsequent regenerative hyperplasia for bladder tumors in rats.

CHEMICAL	CAS NO.	PC CODE		_	QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
				DATE	METHOD	
			Suggestive Evidence Of			
Orthosulfamuron	213464-77-3	108209	Carcinogenic Potential	10/26/2006	RfD Approach	Thyroid tumors in Han Wistar rats (M)
			Likely to be Carcinogenic to			
Oryzalin	19044-88-3	104201	Humans	6/25/2003	Q1* = 7.79 E-3 (3/4)	Thyroid & Skin (M & F) and Mammary (F) tumors in F344 rats
			Likely To Be Carcinogenic To			Liver tumors in F344 rats (M)
Oxadiazon	19666-30-9	109001	Humans	5/1/2001	Q1* = 7.11 E-2 (3/4)	Liver tumors in CD-1 mice (M & F)
			Group CPossible Human			
Oxadixyl	77732-09-3	126701	Carcinogen	1/4/1989	Q1* = 5.3 E-2 (2/3)	Liver tumors in Han-Wistar rats (M & F)
			Group EEvidence of Non-			
Oxamyl	23135-22-0	103801	carcinogenicity for Humans	11/5/1996	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Oxydemeton-methyl	301-12-2	058702	To Humans	7/24/1997	NR	Not Applicable
			Group CPossible Human			
Oxyfluorfen	42874-03-3	111601	Carcinogen		Q1* = 7.32 E-2 (3/4)	Liver tumors in CD-1 mice (M)
			Group DNot Classifiable as to			
Oxytetracycline	2058-46-0	006308	Human Carcinogenicity	12/18/1992	NR	Not Applicable
			Group DNot Classifiable As			
Oxytetracycline	79-57-2	006304	To Human Carcinogenicity	12/18/1992	NR	Not Applicable
			Group DNot Classifiable As			
Oxytetracycline calcium	7179-50-2	006321	To Human Carcinogenicity	12/18/1992	NR	Not Applicable
			Group BProbable Human			Kidney & Liver tumors in F344 rats (M & F)
Oxythioquinox	2439-01-2	054101	Carcinogen		Q1* = 3.42 E-2 (3/4)	Lung tumors in NMRI mice (M)
			Group DNot Classifiable as to			
Paclobutrazol	76738-62-0	125601	Human Carcinogenicity	6/23/1994	NR	Not Applicable
			Not Likely To Be Carcinogenic			Liver tumors in B6C3F1 mice (M & F); Established a mitogenic
Paradichlorobenzene	106-46-7	061501	To Humans	6/5/2007	NR	mode of action for liver tumors.
			Group DNot Classifiable as to			
Paranitrophenol	100-02-7	056301	Human Carcinogenicity	5/14/1996	NR	Not Applicable
			Group EEvidence of Non-			
Paraquat dichloride	1910-42-5	061601	carcinogenicity for Humans	4/19/2000	NR	Not Applicable
						Adrenal, Thyroid & Pancreas tumors in Osborne-Mendel rats
			Group CPossible Human			(M)
Parathion, ethyl-	56-38-2	057501	Carcinogen	9/11/1991	RfD Approach	Pancreas tumors in Wistar rats (M)
			Not Likely To Be Carcinogenic			` '
Pebulate	1114-71-2	041403	To Humans	12/7/1998	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group CPossible Human	IDAIL	INICITIOD	
Pendimethalin	40487-42-1	108501	Carcinogen	7/24/1992	RfD Approach	Thyroid tumors in Sprague-Dawley rats (M & F)
			Suggestive Evidence of Carcinogenicity, but Not		по предоления	and the second s
			Sufficient to Assess Human			
Penoxulam	219714-96-2	119031	Carcinogenic Potential	3/24/2004	NR	Mononuclear Cell Leukemia in Fisher 344 rats (M)
			Group CPossible Human			
Pentachloronitrobenzene (PCNB)	82-68-8	056502	Carcinogen	12/18/1992	RfD Approach	Thyroid tumors in CD rats (M)
			Group BProbable Human			Liver & Vascular (M & F) and Adrenal (M) tumors in B6C3F1
Pentachlorophenol	87-86-5	063001	Carcinogen	1/3/1991	Not Determined	mice
			Likely to be Carcinogenic to			
Permethrin	52645-53-1	109701	Humans	10/23/2002	Q1* = 9.567 E-3 (3/4)	Lung (F) & Liver (M & F) tumors in CD-1 mice
			Group DNot Classifiable as to			
Phenmedipham	13684-63-4	098701	Human Carcinogenicity	4/28/1993	NR	Not Applicable
			Suggestive Evidence of			Vascular tumors in Wistar rats (F) & C5B1/10JfCD-1/Alpk mice (M & F) following oral exposure.
			Carcinogenicity, but Not			
			Sufficient to Assess Human			Vascular tumors in Alderley Park mice (F) following dermal
PHMB	32289-58-0	111801	Carcinogenic Potential	4/9/2003	NR	exposure
			Group EEvidence of Non-			
Phorate	298-02-2	057201	carcinogenicity for Humans	12/30/1993	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Phosalone	2310-17-0	097701	To Humans	8/12/1999	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Phosmet	732-11-6	059201	Carcinogenic Potential	10/27/1999	NR	Liver (M & F) & Mammary (F) tumors in B6C3F1 mice
			Group CPossible Human			
Phosphamidon	13171-21-6	018201	Carcinogen	5/31/1989	NR	Bladder & Liver tumors in Sprague-Dawley rats (M)
<u> </u>			Group EEvidence of Non-			
Phostebupirim	96182-53-5	129086	carcinogenicity for Humans	4/27/1993	NR	Not Applicable
·			Group EEvidence of Non-			
Picloram Acid	1918-02-1	005101	carcinogenicity for Humans	4/1/1994	NR	Not Applicable
			Group EEvidence of Non-			
Picloram Acid Ethylhexyl Ester	2545-60-0	005103	carcinogenicity for Humans	4/1/1994	NR	Not Applicable
			Group EEvidence of Non-			
Picloram Acid Potassium Salt	35832-11-2	005104	carcinogenicity for Humans	4/1/1994	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
Picloram Acid			Group EEvidence of Non-	DATE	METHOD	
Triisopropanolamine Salt	6753-47-5	005102	carcinogenicity for Humans	4/1/1994	ND	Not Applicable
Thisopropariolarnine Sait	0733-47-3	003102	Data Are Inadequate for an	4/1/1994	INIX	Not Applicable
			Assessment of Human			
Pinoxaden	293973-20-8	147500	Carcinogenic Potential	5/18/2005	ND	Not Applicable
I IIIOAAdeII	293973-20-0	147300	Group CPossible Human	3/10/2003	RfD and MOE	Not Applicable
Piperonyl butoxide	51-03-6	067501	Carcinogen	6/7/1005	Approaches	Liver tumors in CD-1 mice (M & F)
i iperoriyi butoxide	31-03-0	007301	Carcinogen	0/1/1993	Арргоаспез	Tumors at multiple sites (Liver and Lung in M & F; Ovary and
			Likely to be Carcinogenic to			Mammary in F) in Swiss mice
Pirimicarb	23103-98-2	106101	Humans	7/13/2005	O1* - 3 526 F -2 (3/4)	Lung tumors in CD-1 mice (F)
Pirimiphos-methyl	29232-93-7	108101	Cannot Be Determined	1/29/1998		Not Applicable
Filliniphos-metriyi	29232-93-1	106102	Cannot be Determined	1/29/1990	INIX	Not Applicable
			Inadequate Information to			
Polymeric Betaine		103679	Assess Carcinogenic Potential	10/3/2006	ND	Not Applicable
Polyment betaine		103679		10/3/2006	INIX	Not Applicable
Potassium dichromate	7778-50-9	068302	Not Likely To Be Carcinogenic To Humans	8/28/2001	ND	Not Applicable
Potassium dichromate	7776-50-9	000302		0/20/2001	INIX	Not Applicable
Potassium Mefluidide	83601-83-6	114003	Not Likely To Be Carcinogenic To Humans	5/30/2007	ND	Net Appliechie
Potassium Menuidide	03001-03-0	114003	Not Likely To Be Carcinogenic	5/30/2007	INIX	Not Applicable
Prallethrin	23031-36-9	128722	To Humans	6/27/2003	ND	Not Applicable
Pralleumin	23031-30-9	120122	Group DNot Classifiable as to		INIX	Not Applicable
Deire in alfance on seathard	00000 54 0	100070			ND	Net Appliechie
Primisulfuron-methyl	86209-51-0	128973	Human Carcinogenicity Group CPossible Human	5/3/1990	INK	Not Applicable
Dan alda an	07747.00.5	400054		7/4/4000	04* 4554 (0/0)	Lives toward in CD Assiss (M. 9. E)
Prochloraz	67747-09-5	128851	Carcinogen	7/1/1988	Q1* = 1.5 E-1 (2/3)	Liver tumors in CD-1mice (M & F)
Dan sa mai da na a	20000 40 0	100011	Group BProbable Human	4/5/4004	04* 4 000 5 0 (0(4)	Testes & Pituitary tumors in Osborne-Mendel rats (M & F)
Procymidone	32809-16-8	129044	Carcinogen	4/5/1991	Q1* = 1.339 E-2 (3/4)	Liver tumors in B6C3F1 mice (F)
Des districts	00004 04 0	440004	Group CPossible Human	0/40/4004	D(D A	Thyroid & Pancreas tumors in Sprague- Dawley rats (M & F)
Prodiamine	29091-21-2	110201	Carcinogen	6/10/1991	RfD Approach	Fibrosarcomas in CD-1 mice (M)
D. G. G. G.	44400 00 7	444404	Group EEvidence of Non-	0/0/4005	ND	Nich Acceptable
Profenofos	41198-08-7	111401	carcinogenicity for Humans	2/6/1995	NK	Not Applicable
B. J. S. F. S.	407077 50 0	110000	Not Likely To Be Carcinogenic	4/4 4/0000	LID.	No. A. P. of I.
Prohexadione	127277-53-6	112600	To Humans	4/14/2000	NK	Not Applicable
L .	1010 10 5	200004	Group DNot Classifiable as to			
Prometon	1610-18-0	080804	Human Carcinogenicity	11/25/1992	NK	Not Applicable
<u></u>	7007.40.6	202225	Group EEvidence of Non-	7/00/465		
Prometryn	7287-19-6	080805	carcinogenicity for Humans	7/26/1994	NK	Not Applicable

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION	REPORT DATE	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
Pronamide	23950-58-5	101701	Group BProbable Human Carcinogen	12/10/2001	Q1* = 2.59 E-2 (3/4)	Testes (M) & Thyroid (M & F) tumors in Crl:CD(SD)BR rats Liver tumors B6C3F1 mice (M)
Propachlor	1918-16-7	019101	Likely to be Carcinogenic to Humans	10/16/1997	Q1* = 3.2 E-2 (3/4)	Stomach (M) tumors in Fischer 344 rats Thyroid (M & F) & Ovary (F) tumors in Sprague-Dawley rats Liver tumors in CD-1 mice (M)
Propamocarb hydrochloride	25606-41-1	119302	Not Likely To Be Carcinogenic To Humans	5/31/2000	NR	Not Applicable
Propanil	709-98-8	028201	Suggestive Evidence of Carcinogenicity, but Not Sufficient to Assess Human Carcinogenic Potential	6/19/2001	NR	Testes & Liver tumors in Sprague-Dawley rats (M)
Propargite	2312-35-8	097601	Group BProbable Human Carcinogen	7/23/1992	Q1* = 1.92 E-1 (3/4)	Jejunum tumors in Crl:CDBR rat (M & F)
Propazine	139-40-2	080808	Not Likely To Be Carcinogenic To Humans	12/8/2005	NR	Mammary tumors in Sprague Dawley rats (F); Established a neuroendocrine mode of action for mammary tumors in rats.
Propetamphos	31218-83-4	113601	Not Likely To Be Carcinogenic To Humans	10/31/1998	NR	Not Applicable
Propiconazole	60207-90-1	122101	Group CPossible Human Carcinogen	9/14/1992	RfD Approach	Liver tumors in CD-1 mice (M)
Propoxur	114-26-1	047802	Group BProbable Human Carcinogen	6/17/1996	Q1* = 3.69 E-3 (3/4)	Bladder tumors in Wistar rats (M & F) Liver tumors in B6C3F1 mice (M)
Propoxycarbazone-Sodium	181274-15-7	122019	Not Likely To Be Carcinogenic To Humans	4/6/2004	NR	Not Applicable
Prosulfuron	94125-34-5	129031	Data Are Inadequate for an Assessment of Human Carcinogenic Potential	1/24/2000	NR	Not Applicable
Prothioconazole	178928-70-6	113961	Not Likely To Be Carcinogenic To Humans	12/31/2007	NR	Not Applicable
Pymetrozine	123312-89-0	101103	Likely to be Carcinogenic to Humans	9/22/1999	Q1* = 1.19 E-2 (3/4)	Liver tumors in Tif:RAIf(SPF) Sprague-Dawley rats (F) Liver tumors in Tif:MAGf(SPF) mice (M & F)
Pyraclostrobin	175013-18-0	099100	Not Likely To Be Carcinogenic To Humans	2/15/2007	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION	REPORT DATE	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Likely to be Carcinogenic to			
Pyraflufen ethyl	129630-19-9	030090	Humans	10/8/2002	Q1* = 3.32 E-2 (3/4)	Liver tumors in (SPF) ICR Crj CD-1 mice (M &F)
			Suggestive Evidence of			Eye tumors in Wistar rats (M)
Pyrasulfatole	365400-11-9	000692	Carcinogenic Potential	5/17/2007	NR	Urinary bladder tumors in C57BL mice (M & F)
			Not Likely To Be Carcinogenic			
Pyrazon	1698-60-8	069601	To Humans	7/28/2005	NR	Not Applicable
Pyrethrins	8003-34-7	069001	Not Likely To Be Carcinogenic To Humans at doses that do not cause mitogenic repsonse in the liver cell proliferation	2/14/2008	NR	Liver tumors in Crl:CD® (SD)IGS BR rats (F); Established a non-genotoxic mitogenic mode of action for liver tumors.
			Group EEvidence of Non-			
Pyridaben	96489-71-3	129105	carcinogenicity for Humans	5/11/1994	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Pyridalyl	179101-81-6	295149	To Humans	8/26/2004	NR	Not Applicable
Pyridate	55512-33-9	128834	Not Likely To Be Carcinogenic To Humans	1/24/2000	NR	Not Applicable
Pyrimethanil	53112-28-0	288201	Group CPossible Human Carcinogen	2/11/1997	MOE Approach	Thyroid tumors in Sprague-Dawley rats (M &F)
Pyriproxyfen	95737-68-1	129032	Group EEvidence of Non-carcinogenicity for Humans	8/15/1995	NR	Not Applicable
			Group CPossible Human			Kidney tumors in Crl:CDBR rats (M)
Pyrithiobac-sodium	123343-16-8	078905	Carcinogen	9/5/1995	Q1* = 1.05 E-3 (3/4)	Liver tumors in CD-1 mice (M)
Pyroxsulam	422556-08-9	108702	Not Likely To Be Carcinogenic To Humans	7/12/2007	NR	Not Applicable
Quinchlorac	84087-01-4	128974	Group DNot Classifiable as to Human Carcinogenicity	8/26/1992	NR	Not Applicable
Quinoxyfen	124495-18-7	055459	Not Likely To Be Carcinogenic To Humans	1/28/2003	NR	Not Applicable
Quizalofop ethyl	76578-14-8	128711	Group DNot Classifiable as to Human Carcinogenicity	3/17/1988	NR	Not Applicable
Quizalolop outly!	70070 1-70	120711	Group DNot Classifiable as to		1111	Trock applicable
Quizalofop-P ethyl	100646-51-3	128709	Human Carcinogenicity	8/10/2006	NR	Not Applicable
. ,			Likely to be Carcinogenic to			Liver tumors in Sprague-Dawley rats (F)
Resmethrin	10453-86-8	097801	Humans	5/25/2005	Q1* = 5.621 E-2 (3/4)	Liver tumors in Swiss Mice (M)
Dimoulfuron	122024 40 0	120000	Not Likely To Be Carcinogenic	2/40/4000	ND	Not Applicable
Rimsulfuron	122931-48-0	129009	To Humans	2/19/1998	INK	Not Applicable

CAS NO.	PC CODE				TUMOR SITES/ STRAIN/ SPECIES/ SEX
		Group EEvidence of Non-			
83-79-4	071003	carcinogenicity for Humans	10/5/1988	NR	Not Applicable
		Not Likely To Be Careinagenia			
372137-35-4	118203		7/22/2000	NR	Not Applicable
372137-33-4	110203		1/22/2009	INIX	Not Applicable
28434-00-6	004004		12/2/2003	NR	Kidney tumors in Sprague-Dawley rats (M)
	00.00.		, _ , _ 5 6 6		ruaney tamere in opragae Damey rate (iii)
74051-80-2	121001	To Humans	3/19/2003	NR	Not Applicable
					Mammary tumors in Sprague-Dawley rats (F); Established a
		Not Likely To Be Carcinogenic			mode of action for neuroendocrine disruption for mammary
122-34-9	080807	To Humans	4/14/2005	NR	tumors in rats.
		Group CPossible Human			
87392-12-9	108800	Carcinogen	9/28/2001	MOE Approach	Liver tumors in Charles River CD (SD)BR rats (F)
50723-80-3	103901		3/8/2006	NR	Not Applicable
					Oral Mucosa and Tongue Mouse B6C3F1 (M & F)
10588-01-9	068304		7/1/2009	Q1* = 7.91 x 10-1	Small Intestines Rat F344 (M & F); Mutagenesis
25155-30-0	079010		7/19/2006		Not Applicable
7775-19-1	011104			NR	Not Applicable
45000 50 0	200004				
15922-78-8	088004		5/16/1995	NR	Not Applicable
4000 40 4	044440		44/04/4000	ND	Not Applicable
1330-43-4	011112		11/24/1993	NK	Not Applicable
12170 04 2	011110		11/24/1002	ND	Not Applicable
12179-04-3	011110	Carcinogenicity For Humans	11/24/1993	INIX	Not Applicable
197166-40-1 ±		Not Likely To Be Carcinogenic			
	110008		9/20/2007	NR	Not Applicable
107 100-10-0	1 10000	1	3/20/2007	INIX	Ποτηφριισασίο
131929-60-7	110003		7/18/2002	NR	Not Applicable
101020 00 7	. 10000		771072002	1111	Testes (M) & Uterine (F) tumors in Wistar rats
148477-71-8	124871		6/10/2004	Q1* = 1.49 F-2 (3/4)	Liver tumors in CD-1 mice (M & F)
	83-79-4 372137-35-4 28434-00-6 74051-80-2 122-34-9	83-79-4 071003  372137-35-4 118203  28434-00-6 004004  74051-80-2 121001  122-34-9 080807  87392-12-9 108800  50723-80-3 103901  10588-01-9 068304  25155-30-0 079010  7775-19-1 011104  15922-78-8 088004  1330-43-4 011112  12179-04-3 011110  187166-40-1 + 187166-15-0 110008  131929-60-7 110003	Group EEvidence of Non-carcinogenicity for Humans	Sarapa	Group EEvidence of Non-   Carcinogenicity for Humans   10/5/1988 NR

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION	TUMOR SITES/ STRAIN/ SPECIES/ SEX
				DATE	METHOD	
			Not Likely To Be Carcinogenic			
Spiromesifen	283594-90-1	024875	To Humans	5/21/2008	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Spirotetramat	203313-25-1	392201	To Humans	3/26/2009	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Spiroxamine	118134-30-8	120759	To Humans	11/14/2003	NR	Not Applicable
			Group EEvidence of Non-			
Sulfentrazone	122836-35-5	129081	carcinogenicity for Humans	5/7/1996	NR	Not Applicable
			Group EEvidence of Non-			
Sulfosate	81591-81-3	128501	carcinogenicity for Humans	7/26/1994	NR	Not Applicable
			Not Likely to be Carcinogenic			Urinary bladder tumors in Sprague-Dawley rats (F)
Sulfosulfuron	141776-32-1	085601	to Humans	12/16/2008	NR	Urinary bladder (M) and Kidney (M & F) tumors in CD-1 mice
			Not Likely To Be Carcinogenic			
Sulfuryl fluoride	2699-79-8	078003	To Humans	5/24/2001	NR	Not Applicable
			Group EEvidence of Non-			
Sulprofos	35400-43-2	111501	carcinogenicity for Humans	3/26/1996	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Sumithrin	26002-80-2	069005	To Humans	5/30/2006	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Tau-fluvalinate	102851-06-9	109302	To Humans	9/29/2005	NR	Not Applicable
			Group CPossible Human			
TCMTB (Busan 72)	21564-17-0	035603	Carcinogen	8/28/1996	RfD Approach	Testes (M) & Thyroid (F) tumors in Sprague-Dawley rats
			Group CPossible Human			
Tebuconazole	107534-96-3	128997	Carcinogen	9/15/1993	RfD Approach	Liver tumors in NMRI mice (M & F)
			Group EEvidence of Non-			
Tebufenozide	112410-23-8	129026	carcinogenicity for Humans	8/29/1994	NR	Not Applicable
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Tebufenpyrad	119168-77-3	090102	Carcinogenic Potential	7/15/2002	NR	Liver tumors in F344 rats (M & F)
			Group DNot Classifiable as to			
Tebuthiuron	34014-18-1	105501	Human Carcinogenicity	3/1/1993	NR	Not Applicable
						Tumors at multiple sites (Forestomach, Liver, Mammary,
			Group BProbable Human		Q1* = 1.3 E-5 (3/4)	Thyroid, Adrenal, Urinary, Lung) in Fischer 344 rats & B6C3F1
Telone	542-75-6	029001	Carcinogen	3/19/2002	(Inhalation)	mice (M & F)

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Suggestive Evidence of			
Tembotrione	335104-84-2	012801	Carcinogenic Potential	5/22/2007	RfD Approach	Eye tumors in Wistar rats (M)
			Data Are Inadequate for an			
			Assessment of Human			
Tepraloxydim	149979-41-9	121005	Carcinogenic Potential	2/27/2001	NR	Not Applicable
			Group EEvidence of Non-			
Terbacil	5902-51-2	012701	carcinogenicity for Humans	9/30/1994	NR	Not Applicable
			Group EEvidence of Non-			
Terbufos	13071-79-9	105001	carcinogenicity for Humans	3/9/1994	NR	Not Applicable
			Group DNot Classifiable as to			
Terbuthylazine	5915-41-3	080814	Human Carcinogenicity	8/24/1994	NR	Not Applicable
			Group CPossible Human			Tumors at multiple sites (Mammary, Liver, Thyroid, Testes in
Terbutryn	886-50-0	080813	Carcinogen	3/3/1988	NR	CD rats (M & F)
			Group BProbable Human			Tumors at multiple sites (Liver, Bile duct, Mammary,
Terrazole	2593-15-9	084701	Carcinogen	6/29/1999	Q1* = 3.33 E-2 (3/4)	Thyroid,Testes) in Sprague-Dawley rats (M & F)
			Likely to be Carcinogenic to		<u> </u>	Adrenal & Thyroid tumors in Sprague-Dawley rats (M)
Tetrachlorvinphos	961-11-5	083701	Humans	3/7/2002	Q1* = 1.83 E-3 (3/4)	Liver tumors; B6C3F1 mice (F)
			Likely to be Carcinogenic to			
Tetraconazole	112281-77-3	120603	Humans	1/11/2000	Q1* = 2.3 E-2 (3/4)	Liver tumors in Crl:CD-1 (ICR) mice (M &F)
Tetramethrin	7696-12-0	069003	Group CPossible Human Carcinogen	12/11/1989	NR	Testes tumors in CR CD-1 rats, Sprague-Dawley rats & Long- Evans Hooded rats (M)
Thiabendazole	148-79-8	060101	Multiple Descriptors: Likely to be Carcinogenic to Humans at High Does; Not Likely to be Carcinogenic to Humans at Low Doses	3/8/2002	MOE Approach	Thyroid tumors in Sprague-Dawley Crl:CD BR rats (M & F); Established a hormonal mode of action for thyroid tumors.
			Likely to be Carcinogenic to			Thyroid (M & F) & Uterine (F) tumors in Wistar rats
Thiacloprid	111988-49-9	014019	Humans	3/26/2003	Q1* = 4.06 E-2 (3/4)	Ovarian tumors in B6C3F mice (F)
Thiamethoxam	153719-23-4	060109	Not Likely To Be Carcinogenic To Humans	6/13/2005	NR	cytotoxic, regenerative proliferative, non-genotoxic mode of action for liver tumors in mice.
			Suggestive Evidence Of			
Thiazopyr (MON 13200)	117718-60-2	129100	Carcinogenic Potential	12/6/2007	NR	Kidney tumors in Sprague Dawley rats (M & F) )
			Not Likely To Be Carcinogenic			
Thidiazuron	51707-55-2	120301	To Humans	8/31/2005	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Not Likely To Be Carcinogenic			
			To Humans at doses that do			Urinary bladder tumors in C57BL/6J mice (M &F); Established
			not cause urothelium		 	a cytotoxicity and regeneration proliferation mode of action for
Thiencarbazone methyl	317815-83-1	015804	cytotoxicity	2/29/2008	NR	urinary bladder tumors in mice.
			Not Likely To Be Carcinogenic			
Thifensulfuron methyl	79277-67-1	128845	To Humans	12/12/2006	NR	Not Applicable
T	20040 77 0	100101	Group DNot Classifiable as to			
Thiobencarb (Bolero)	28249-77-6	108401	Human Carcinogenicity	6/10/1996	NR	Not Applicable
This is a share to the same of the	04005.00.4	400000	Group DNot Classifiable as to		ND	No. Co. Production
Thiocyclam hydrogen oxalate	31895-22-4	128868	Human Carcinogenicity	9/15/1994	NK	Not Applicable
This discul-	50000 00 0	444504	Group BProbable Human	0/40/4000	MOE Ammanah	Testes tumors in Sprague-Dawley rat (M)
Thiodicarb	59669-26-0	114501	Carcinogen	6/10/1996	MOE Approach	Liver tumors in CD-1 mice (M & F)
			Likely to be Carcinogenic to			Thyroid tumors in F344 rats (M &F)
Thiophanate-methyl	23564-05-8	102001	Humans	8/24/1999	Q1* = 1.16 E-2 (3/4)	Liver tumors in CD-1 mice (M & F)
			Not Likely To Be Carcinogenic			
Thiram	137-26-8	079801	To Humans	4/14/2003	NR	Not Applicable
			Likely to be Carcinogenic to			
Tolyfluanid	731-27-1	309200	Humans	6/18/2002	Q1* = 1.59 E-3 (3/4)	Thyroid tumors in Wistar rats (M & F)
Topramezone	210631-68-8	123009	Multiple Descriptors: Not Likely to be Carcinogenic to Humans at Doses that Do Not Alter Rat Thyroid Hormone Homeostasis	5/19/2005	NR	Thyroid tumors in Wistar rats (M & F); Established a hormonal mode of action for thyroid tumors observed only at an excessive dose.
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			Testicular tumors in Wistar rats (M)
Tralkoxydim	87820-88-0	121000	Carcinogenic Potential	6/30/2004	NR	Ovarian tumors in Syrian Golden hamsters (F)
			Group CPossible Human			Thyroid tumors in Wistar rats (M)
Triadimefon	43121-43-3	109901	Carcinogen	12/4/1996	RfD Approach	Liver tumors in NMRI mice (M & F)
			Group CPossible Human			
Triadimenol	55219-65-3	127201	Carcinogen	1/29/1988	NR	Liver tumors in CF1/W74 mice (F)
			Group CPossible Human			Kidney tumors in Sprague-Dawley rats (M)
Triallate	2303-17-5	078802	Carcinogen	1/12/1994	Q1* = 7.17 E-2 (3/4)	Liver tumors in B6C3F1 mice (F)
			Group EEvidence of Non-		. ,	
Triasulfuron	82097-50-5	128969	carcinogenicity for Humans	2/27/1991	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Triazamate	112143-82-5	128100	To Humans	12/1/1997	NR	Not Applicable

CHEMICAL	CAS NO.	PC CODE			QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group CPossible Human	IDATE	IMETHOD	
Tribenuron methyl	101200-48-0	128887	Carcinogen	7/14/1989	NR	Mammary tumors in Sprague-Dawley rats (F)
The circuit in circuity.	101200 100		Multiple Descriptors: Likely to	.,,		manmary tamero in opragao 2 amoj rato (i )
			be Carcinogenic to Humans			
			(High Doses); Not Likely to be			
			Carcinogenic to Humans (Low			Liver (M), Lung (F), & Small intestine (M & F) tumors in
Tribufos	78-48-8	074801	Doses)	5/22/1997	MOE Approach.	CD-1 mice
11100100	70 10 0	07 100 1	, and the second	0,22,1001	MOL Approudin	
T 11	4.4075 57.4	000440	Group DNot Classifiable As	0/04/000	N.D.	
Tributyltin maleate	14275-57-1	083118	To Human Carcinogenicity	3/31/2005	NR	
			Multiple Descriptors: Likely to			
			be Carcinogenic to Humans			
			(High Doses), Not Likely to be			
			Carcinogenic to Humans (Low			Kidney & Lung tumors in Fischer 344 rats (M & F)
Trichlorfon	52-68-6	057901	Doses)	7/15/1999	NR	Mammary tumors in CD-1 mice (F)
			Group DNot Classifiable as to			
Triclopyr	55335-06-3	116001	Human Carcinogenicity	5/9/1996	NR	Not Applicable
			Not Likely To Be Carcinogenic			Liver tumors in CD-1 mice (M & F); Established a PPARa
Triclosan	3380-34-5	054901	To Humans	1/4/2008	NR	mode of action for liver tumors.
			Group CPossible Human			
Tridiphane	58138-08-2	123901	Carcinogen	4/22/1986	NR	Liver tumors in B6C3F1 mice (F)
			Not Likely To Be Carcinogenic			
Trifloxystrobin	141517-21-7	129112	To Humans	6/16/1999	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Trifloxysulfuron	290332-10-4	119009	To Humans	7/22/2003	NR	Not Applicable
			Group EEvidence of Non-			
Triflumizole	68694-11-1	128879	carcinogenicity for Humans	8/10/1993	NR	Not Applicable
			Group CPossible Human			Thyroid, Renal pelvis & Urinary bladder tumors in Fischer 344
Trifluralin	1582-09-8	036101	Carcinogen	4/11/1986	Q1* = 2.93 E-3 (3/4)	rats (F)
			Group CPossible Human			
Triflusulfuron-methyl	126535-15-7	129002	Carcinogen	5/28/1996	RfD Approach	Testes tumors in CD-1 rats (M)
			Suggestive Evidence of			
			Carcinogenicity, but Not			
			Sufficient to Assess Human			
Triforine	26644-46-2	107901	Carcinogenic Potential	6/29/2004	NR	Liver (M) & Lung (F) tumors in Crl:CD-1 mice
			Not Likely To Be Carcinogenic			
Trinexapac-Ethyl	95266-40-3	112602	To Humans	9/5/2008		

CHEMICAL	CAS NO.	PC CODE	CANCER CLASSIFICATION	_	QUANTIFICATION METHOD	TUMOR SITES/ STRAIN/ SPECIES/ SEX
			Group BProbable Human			Pituitary & Leydig cell tumors in Wistar rats (M &F)
Triphenyltin hydroxide (TPTH)	76-87-9	083601	Carcinogen	5/24/1990	Q1* = 1.83 E-0 (3/4)	Liver tumors in NMRI mice (M &F)
			Not Likely To Be Carcinogenic			
Triticonazole	131983-72-7	125620	To Humans	6/15/2006	NR	Not Applicable
			Not Likely To Be Carcinogenic			
Troysan polyphase (IPBC)	55406-53-6	107801	To Humans	12/4/1996	NR	Not Applicable
			Group BProbable Human			Multiple tumors (Lung, Blood vessels, Liver, Kidney) in multiple
UDMH	57-14-7	600018	Carcinogen	7/26/1991	Q1* = 4.6 E-1 (2/3)	species, strains & studies.
			Group EEvidence of Non-			
UMP-488 (PAL 6000)	111578-32-6	129025	carcinogenicity for Humans	5/6/1994	NR	Not Applicable
			Group CPossible Human			
Uniconazole	83657-22-1	128976	Carcinogen	10/11/1990	NR	Liver tumors in CD-1 mice (M)
			Group CPossible Human	2/22/222		
Vinclozolin	50471-44-8	113201	Carcinogen	6/20/2000	MOE Approach	Leydig cell tumors in Wistar rats (M)
7.0	50045.07.0	400004	Group CPossible Human	0/07/4000		
Zeta-Cypermethrin	52315-07-8	129064	Carcinogen	9/27/1988	NK	Lung tumors in Alderly Park SPF Swiss strain mice (F)
			Suggestive Evidence of			
			Carcinogenicity, but Not			
7:	407.00.4	00.4005	Sufficient to Assess Human	0/0/0000	ND	Hemangiomas in CD(SD)BR rats (M)
Ziram	137-30-4	034805	Carcinogenic Potential	2/6/2003	INK	Preputial gland tumors in F344 rats (M)
7	450050 00 5	404700	Not Likely To Be Carcinogenic	0/7/0004	ND	Not Applicable
Zoxamide	156052-68-5	101702	To Humans	2/7/2001	INK	Not Applicable